

Ndufc2 Gene Inhibition Is Associated With Mitochondrial Dysfunction and Increased Stroke Susceptibility in an Animal Model of Complex Human Disease

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Background—The genetic basis of stroke susceptibility remains to be elucidated. STR1 quantitative trait locus (STR1/QTL) was identified on rat chromosome 1 of stroke-prone spontaneously hypertensive rat (SHRSP) upon Japanese-style stroke-permissive diet (JD), and it contributes to 20% of the stroke phenotype variance.

Methods and Results—Nine hundred eighty-six probe sets mapping on STR1 were selected from the Rat RAE230A array and screened through a microarray differential expression analysis in brains of SHRSP and stroke-resistant SHR (SHRSR) fed with either regular diet or JD. The gene encoding *Ndufc2* (NADH dehydrogenase [ubiquinone] 1 subunit), mapping 8 Mb apart from STR1/QTL Lod score peak, was found significantly down-regulated under JD in SHRSP compared to SHRSR. *Ndufc2* disruption altered complex I assembly and activity, reduced mitochondrial membrane potential and ATP levels, and increased reactive oxygen species production and inflammation both in vitro and in vivo. SHRSR carrying heterozygous *Ndufc2* deletion showed renal abnormalities and stroke occurrence under JD, similarly to SHRSP. In humans, T allele variant at *NDUFC2*/rs11237379 was associated with significant reduction in gene expression and with increased occurrence of early-onset ischemic stroke by recessive mode of transmission (odds ratio [OR], 1.39; CI, 1.07–1.80; $P=0.012$). Subjects carrying TT/rs11237379 and A allele variant at *NDUFC2*/rs641836 had further increased risk of stroke (OR=1.56; CI, 1.14–2.13; $P=0.006$).

Conclusions—A significant reduction of *Ndufc2* expression causes complex I dysfunction and contributes to stroke susceptibility in SHRSP. Moreover, our current evidence may suggest that *Ndufc2* can contribute to an increased occurrence of early-onset ischemic stroke in humans. (*J Am Heart Assoc.* 2016;5:e002701 doi: 10.1161/JAHA.115.002701)

Key Words: complex I • early-onset ischemic stroke • knockout rat model • mitochondria • *Ndufc2* • stroke-prone spontaneously hypertensive rat

It is known that both environmental and genetic factors contribute to increased stroke predisposition in humans.^{1,2} Unraveling the genetic causes of stroke may take significant advantage of the availability of an inbred animal

model, the stroke-prone spontaneously hypertensive rat (SHRSP).³ The latter represents a suitable model for studies on the human disease⁴ and, based on previous investigations, carries at least 3 quantitative trait loci (QTL) for stroke

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susceptibility.⁵ One of them (STR1) maps on rat chromosome 1; it is included between *Klk* and *Mt1pa* markers, extending for an overall 80 cM length, and shows a 6.7 Lod score peak in correspondence of the anonymous marker, *D1Mit3*. STR1 contributes to 20% of overall stroke phenotype variance.⁵ Notably, its effect was confirmed by congenic experimentation.⁶ In fact, congenic animals carrying SHRSP/STR1 within the stroke-resistant (SHRSR) genomic background had increased stroke occurrence. However, no apparent suitable candidate genes were initially discovered near the Lod score peak area. In the attempt to identify the gene(s) responsible for stroke phenotype within STR1, we performed, based on previous literature,⁷ a microarray differential expression analysis of all sequences contained inside STR1 in brains of both SHRSP and SHRSR fed with either regular or stroke permissive diet. One gene was found to be differentially modulated (ie, down-regulated) in brains of SHRSP. This gene was *Ndufc2* (NADH dehydrogenase [ubiquinone] 1, subcomplex unknown, 2), mapping 8 Mb apart from the Lod score peak.

Here, we report both in vitro and in vivo evidence of the significant functional impact of *Ndufc2*-reduced expression on mitochondrial function, oxidative stress accumulation, and cell viability. We describe the stroke phenotype of SHRSR carrying heterozygous *Ndufc2* deletion. Moreover, we report results of a genetic screening carried out in a human cohort of subjects with early-onset ischemic stroke by testing available NDUFC2 single nucleotide polymorphisms (SNPs), which may generate the hypothesis that *Ndufc2* is a genetic risk factor for stroke also in humans.

Methods

Brain Expression Profiling of Potential Candidate Genes Within STR1/QTL on Rat Chromosome 1 in SHRSP

Ten 6-week-old SHR/molBbb (SHRSR) and ten 6-week-old SHRSP/Bbb (SHRSP) rats, both inbred for 20 years at the MDC center in Berlin and derived from the original colonies established in Japan, were fed with either regular (RD; n=5) or Japanese-style diet (JD; n=5) for 4 weeks and were then sacrificed. Brains were removed, immediately frozen, and subsequently extracted for total RNA by using the QiAGEN's RNeasy Total RNA Isolation kit (Qiagen GmbH, Hilden, Germany). Preparation quality was assessed by agarose-formaldehyde gel electrophoresis. For synthesis of double-stranded cDNA from 15 µg of total RNA, the cDNA Synthesis System kit (Roche Diagnostics, Indianapolis, IN) was used. Biotinylated cRNA was synthesized with PerkinElmer's (Waltham, MA) nucleotide analogs using Ambion's MEGAScript T7 kit (Ambion, Austin, TX). After fragmenting

the cRNA for target preparation using the standard Affymetrix protocol, 15 µg of fragmented cRNA was hybridized for 16 hours at 45°C to Rat Genome 230 A Array. After hybridization, the arrays were washed and stained with streptavidin-phycoerythrin in the Affymetrix Fluidics Station 400 and further scanned using the AFFYMETRIX GeneChip Scanner 3000 7G (Affymetrix Inc, Santa Clara, CA). Image data were analyzed with GCOS 1.4 using Affymetrix default analysis settings and global scaling as a normalization method. All arrays passed our formal quality criteria (3'-5' ratio of Gapdh <3; RawQ <3) and were quantile-normalized using Robust Multiarray Average (RMA).⁸ Outliers in replicates were removed according to a Nalimov test *P*-value threshold of 10⁻³.⁹ After removal of probe sets located outside the major stroke QTL interval on rat chromosome 1 (92–220 Mbps), the remaining 986 probe sets were analyzed using ANOVA statistic followed by a false discovery rate (FDR) multiple testing correction.¹⁰ Probes that underwent 0.1% FDR (N=87) were k-means clustered. The cluster analysis was done after applying probe-set standardization using an average Euclidean distance function. *K* was selected according to the absolute minimum in the Davies Bouldin cluster estimation procedure.¹¹ Furthermore, to estimate contribution of treatment and strain differences and the interaction to the general ANOVA effect, a 2-way ANOVA was performed on probe sets representing *Ndufc2*.

Data of microarray analysis are available online (<https://www.ebi.ac.uk/arrayexpress/experiments/E-MTAB-1102>; accession numbers: ArrayExpress acc#: E-MTAB-1102).

Sequencing of *Ndufc2* in the 2 Parental Strains

As previously reported,¹² genomes of SHR/NHsd (Harlan Laboratories, Bethesda, MD) and of SHRSP/Gla (Western Infirmary, Glasgow, UK) were sequenced on an Illumina platform to a depth of at least 23× (Illumina, San Diego, CA). In brief, 5 µg of DNA was used to construct paired-end whole-genome libraries with 300- to 600-base-pair (bp) insert size. Genomic DNA was prepared from liver or spleen and quality checked before sequencing on an Illumina HiSeq2000, following the manufacturer's instructions. Quality-filtered Illumina paired-end and/or mate pair reads were mapped to the Brown Norway reference genome, RGSC-3.4,¹³ using the read alignment software Burrows-Wheeler Aligner (BWA-0.5.8c).¹⁴ Genomic variants (single-nucleotide variant and short indels [1–15 bp]) were detected using the Genome Analysis Toolkit (GATK version 1.0.6001).^{15,16} Sequence differences between strains SHR/NHsd and SHRSP/Gla, compared to other rat strains, were accessed using custom Perl scripts.

Determination of *Ndufc2* Expression by Reverse-Transcription Polymerase Chain Reaction in Brain Tissue and in Cellular Extracts

Two micrograms of total RNA were used for cDNA synthesis using Superscript III First-Strand (Invitrogen, Carlsbad, CA) and random examer primers according to manufacturer's instructions. The following oligonucleotides were used: *Ndufc2*: forward 5'-GGCTTGCTACATCGGCTTC-3'; reverse 5'-TGATGGTCCCTCACAGCATA-3'; *β-actin*: forward 5'-AGATG ACCCAGATCATGTTTGA-3'; reverse 5'-ATAGGGACATGCGG AGACCG-3'. Real-time quantitative polymerase chain reaction (RT-PCR) was performed using a 2× SYBR Green PCR Master Mix (Applied Biosystems, Foster City, CA) containing the double-stranded DNA-binding fluorescent probe, Sybr Green, and all necessary components except primers. Quantitative PCR conditions included an initial denaturation step of 94°C/10 minutes followed by 40 cycles of 94°C/15 seconds and 60°C/15 seconds. Standards, samples, and negative controls (no-template) were analyzed in triplicate. Concentrations of mRNA were calculated from serially diluted standard curves simultaneously amplified with the unknown samples and corrected for *β-actin* mRNA levels. Levels of *Ndufc2* mRNA in JD-fed rats were compared to levels of RD-fed rats. Levels of *Ndufc2* mRNA in *Ndufc2*-silenced cells were compared to nonsilenced cells.

Western Blotting of Total Proteins Extracted From Brain Tissue and Cellular Extracts

Brain total proteins were extracted as previously reported for tissues.¹⁷ For cellular protein extraction, cells were washed twice with ice-cold PBS and lysed with lysis buffer. Protein concentrations were determined by the Bradford method.¹⁸ Then, 40 μg of total proteins were separated on 12% SDS-PAGE and transferred to polyvinylidene difluoride (PVDF) membranes (Amersham, Piscataway, NJ). Nonspecific binding sites were blocked with 5% nonfat dried milk for 2 hours at room temperature (RT). Membranes were then incubated overnight with the following primary antibodies: anti-*Ndufc2* (1:200) rabbit polyclonal (Catalog No.: NBP1-59610; Novus Biologicals, Littleton, CO); anti-Nf-κBp65 (nuclear factor kappa B; 1:200; Catalog No.: SC-8008; Santa Cruz Biotechnology, Santa Cruz, CA); anti-SOD2 (superoxide dismutase 2, mitochondrial; 1:200; Catalog No.: S5569; Sigma-Aldrich, Milan, Italy), anti-phospho-MAPKp38 (mitogen-activated protein kinase; 1:200) mouse monoclonal (Catalog No.: SC-7973; Santa Cruz Biotechnology); anti-phospho-JNK1 (c-Jun N-terminal kinase 1; 1:200) mouse monoclonal (Catalog No.: SC-6254; Santa Cruz Biotechnology); anti-MAPKp38 (1:200) rabbit polyclonal (Catalog No.: SC-535; Santa Cruz Biotech-

nology); anti-JNK1 (1:200) rabbit polyclonal (Catalog No.: SC-474; Santa Cruz Biotechnology); anti-IκB (nuclear factor of kappa light polypeptide gene enhancer in B-cells inhibitor; 1:200) rabbit polyclonal (Catalog No.: SC-945; Santa Cruz Biotechnology); anti-c-Jun mouse monoclonal antibody (1:200; Catalog No.: L70B11; Cell Signaling Technology, Milan, Italy); anti-β-actin (1:5000; Catalog No.: A5441; Sigma-Aldrich). Secondary antibodies were: 1:5000 antirabbit (Catalog No.: SC-2004; Santa Cruz Biotechnology) and 1:5000 antimouse (Catalog No.: SC-2005; Santa Cruz Biotechnology).

Signals were revealed with an enhanced chemiluminescence detection system (ECL; Amersham) and visualized by a ChemiDoc XRS+imaging system (Bio-Rad, Richmond, CA). Finally, protein levels were normalized using β-actin levels. Protein bands were scanned and quantified densitometrically.

Detection of Brain-Oxidized Proteins

To detect levels of brain-oxidized proteins from total proteins, the Oxyblot Detection kit (Millipore, Milan, Italy) was used as previously reported.¹⁷ Five selected protein bands from immunoblots were scanned and quantified densitometrically. Experiments were performed in triplicate.

In Vitro *Ndufc2* Silencing in A10 Cells

A vascular smooth muscle cell line (A10) obtained from rat embryonic thoracic aorta (purchased from LGC Promochem, Milan, Italy) was cultured in DMEM containing normal glucose (5.5 mmol/L), 10% FBS, and supplemented with penicillin (100 U/mL)/streptomycin (100 μg/mL) at 37°C in 95% O₂ and 5% CO₂. Cells were plated in 100-mm-diameter dishes (4×10⁵) and 60-mm-diameter dishes (1.5×10⁵), passaged upon reaching confluence with 2 mL of trypsin and used at the 12th passage and 70% confluence. Cells were washed with PBS, and then OPTI-MEM—reduced serum medium (Invitrogen, Carlsbad, CA) was added to the cells. *Ndufc2*-specific siRNA (Mission siRNA; Sigma-Aldrich) and a nucleic acid transferring agent, Lipofectamine 2000 (Invitrogen) were incubated in OPTI-MEM—reduced serum medium for 20 minutes at RT to form a siRNA-Lipofectamine complex. The siRNA-Lipofectamine complex-containing medium was added to cells to a final siRNA concentration of 33 nmol/L. Five hours later, the complex-containing medium was replaced with DMEM supplemented with 10% FBS. Cells transfected with Lipofectamine and no-target siRNA (Sigma-Aldrich) were used as controls. Seventy-two hours later, both silenced and nonsilenced cells were used for all analyses described below. Six experiments were performed.

Determination of Reactive Oxygen Species Production in *Ndufc2*-Silenced A10 Cells

Intracellular production of superoxide was evaluated by dichlorodihydrofluorescein ester (DCHF), as previously reported.¹⁷ For this purpose, A10 cells were incubated in 1 mL of 0.05 mmol/L of PBS solution of the permeable agent, DCHF (Sigma-Aldrich), for 30 minutes. Cells were lysed with 2% Triton-X. Lysates were collected on ice, centrifuged at 13 800 g at 4°C for 10 minutes, and then fluorescence intensity was read. In this assay, oxidative stress is quantified by monitoring intracellular DCF content by a fluorometer (Mithras LB 940; Berthold Technologies GmbH & Co. KG, Bad Wildbad, Germany) with excitation at 495 nm and emission at 525 nm. Fluorescence intensity measurements are expressed as arbitrary units.

Characterization of Inflammatory Pathway in *Ndufc2*-Silenced A10 Cells

RNA extracted from both silenced and nonsilenced cells was analyzed by quantitative RT-PCR using an RT²Profiler PCR array (Rat Inflammation array; SuperArray Bioscience Corporation, Frederick, MD). Procedure and data analysis have been previously described.¹⁹ Results are expressed as relative levels of each gene mRNA under condition of either presence or absence of *Ndufc2* referred to the expression of this gene in control cells (that were chosen to represent 1× expression of each gene). Results were considered significant when mRNA expression was 2.5-fold higher or lower than that of control cells. Experiments were performed in triplicate.

Assessment of Necrosis by Flow Cytometry With Propidium Iodide Staining

Cells were harvested by incubation with 2 mL of trypsin for 5 minutes. After addition of 10% FBS DMEM medium, the suspension was centrifuged at low speed at RT. Each pellet was washed with PBS. After washing, 5×10^5 cells were resuspended in 2 mL of cold PBS and centrifuged at 4°C at 500g for 5 minutes. The supernatant was removed, pellets energetically vortexed, and then gently resuspended in 400 μ L of hypotonic solution of propidium iodide (PI; 50 mg/mL in 0.1% sodium citrate plus 0.1% Triton X-100; Sigma-Aldrich). Cells were incubated in the dark at 4°C for 20 minutes. Then, fluorescence-activated cell sorting (FACS) was performed by a FACS Calibur flow cytometer (BD Biosciences, San Jose CA). For this assay, red fluorescence was measured corresponding to the red color of PI (FL-3 detector). To exclude doublets and cell aggregates from the analysis, a doublet discrimination module was used. Peak width was estimated using the coefficient of variation of the signals within each peak. Cells treated with a high dose of H₂O₂ were used as positive control.

Analysis of Mitochondrial Function

Complex I assembly evaluation

Mitochondria were isolated from whole rat brains using the Mitochondria Isolation Kit for Rodent Tissue (Abcam, Cambridge, MA). Preparation of total mitochondrial extracts for Blue Native (BN)-PAGE (BNG) analysis was performed as described previously.^{20,21} Mitochondrial pellet was resuspended at a protein concentration of ≈ 1 mg/mL in a solubilization buffer containing 50 mmol/L of Bis-Tris (pH 7.0), 1% (v/v) *n*-dodecyl- β -D-maltoside, and 750 mmol/L of ϵ -amino-*N*-caproic acid. The suspension was kept on ice for 1 hour with occasional vortexing and was then clarified by centrifugation at 14 000g for 30 minutes. Then, 0.9 mL of the resulting supernatant, containing both membrane and water-soluble proteins, was mixed with 0.1 mL of concentrated BN-PAGE loading buffer (10×) containing 0.75 mol/L of ϵ -amino-*N*-caproic acid and 3% (w/v) Serva blue G-250. Samples were stored at -20°C until analysis. Protein concentrations were determined by bicinchoninic acid protein assay using BSA as a standard. Intact mitochondrial complexes were then separated by electrophoresis using a minigel system (Bio-Rad Mini-Protean 3) and a 4% to 15% gradient acrylamide gel (Mini-Protean Precast Gels; Bio-Rad). Mitochondrial protein complexes were transferred to a methanol-activated PVDF membrane (Bio-Rad) using the BioRad Mini Trans-blot System. Membranes were then blocked overnight at 4°C in 5% milk/PBS solution, washed for 10 minutes in PBS 0.05% Tween-20, and incubated for 3 hours with the primary antibodies (MitoProfile Total OXPHOS Rodent WB Antibody Cocktail; Abcam) to a final concentration of 6.0 μ g/mL in 1% milk/PBS solution. Membranes were washed in PBS 0.05% Tween-20 and incubated for 1 hour with the secondary antibody at RT (goat anti-mouse, 1:5000, HRP-linked). Rabbit polyclonal Hsp60 was used as loading control (Anti-Hsp60 antibody-Loading Control; Abcam) at a concentration of 1 μ g/mL in 1% milk/PBS solution at 1:3000 dilution. Bands were detected using ECL Luminata Crescendo (Millipore), and images were acquired using the ChemiDoc XRS+imaging system. Experiments were performed in triplicate.

Ndufc2-silenced cells were harvested using trypsin and washed twice with PBS. After removal of residual PBS, cells were immediately treated with the Mitochondria Isolation Kit for Cultured Cells (Abcam). Samples were then treated as reported above for brain tissues.

Complex I activity

Brains were dissected out, weighted, wetted from blood, and placed in ice-cold respiration medium (sucrose 250 mmol/L, MgCl₂ 5 mmol/L, EDTA 1 mmol/L [pH 7.4], HEPES 20 mmol/L [pH 7.4], and KH₂PO₄ 2 mmol/L). Tissue was chopped finely with a pair of scissors and rinsed in new

respiration medium to remove any blood. Disrupted tissue was then homogenized on ice by 20 to 30 passes in a tight-fitting glass/Teflon power-driven Potter-Elvehjem homogenizer in ≈ 25 mL of respiration buffer. Homogenate was centrifuged at 600 rpm for 10 minutes at 4°C. The supernatant, rich with mitochondria, was then centrifuged at 4°C at 12 000 rpm for 10 minutes. The external part of the pellet, which is the cleaner part, was resuspended in a small volume of respiration buffer by a miniglass/Teflon power-driven Potter-Elvehjem. Mitochondrial protein concentrations were determined as stated above.¹⁸

Silenced cells were washed once with cold PBS 1 \times and treated with lysis buffer.

Activity of mitochondrial complex I was analyzed using an ELISA microplate assay kit (Abcam). Capture antibody for complex I was precoated in the wells of the microplate, and samples containing 50 μ g of brain mitochondrial extracts and 40 μ g of cell lysates were added to wells. In this assay, complex I activity is measured by oxidation of NADH to NAD⁺, which leads to increased absorbance at 450 nm. Absorbance was read by a Benchmark microplate reader (Bio-Rad). All tests were done in triplicate.

Mitochondrial membrane potential assessment

Brain mitochondrial membrane potential was determined by a fluorescent probe, TMRE (Abcam). Briefly, TMRE reagent was warmed to room temperature and diluted with PBS 1 \times 0.2% BSA to obtain a final TMRE staining working solution at a concentration of 500 nmol/L. Isolated mitochondria (100 μ g) were incubated with 80 μ L of TMRE staining working solution for 10 minutes at 37°C. Fluorescence was determined using a fluorometer (Mithras LB 940; Berthold) microplate reader at 549 nm excitation/575 nm emission. Data are expressed as fluorescent units per μ g protein. Mitochondrial membrane potential of silenced cells was determined by use of the TMRE Mitochondrial Membrane Potential Assay Kit (Abcam). Fluorescence was determined by a FACS Calibur flow cytometer (FL-2 detector). Data are expressed as fluorescent intensity.

ATP-level measurement

Brain mitochondria (250–500 μ g) were plated in a dark 96-multiwell plate on ice and incubated, always on ice, for 5 minutes with respiration buffer (sucrose 250 mmol/L, MgCl₂ 5 mmol/L, EDTA 1 mmol/L, HEPES 20 mmol/L, KH₂PO₄ 2 mmol/L, ADP 1 mmol/L, D-Luciferin, and Luciferase firefly ready from ATP Determination Kit; Invitrogen, Milan, Italy) in the presence of both glutammate (2.5 mmol/L) and malate (2.5 mmol/L; complex I substrates) or succinate (5 mmol/L; complex II substrate). In this assay, quantitative determination of ATP with recombinant firefly luciferase and its substrate, D-luciferin, is based on luciferase's requirement

for ATP in producing light (emission ≈ 560 nm at pH 7.8) from the reaction. Absorbance was quantified by a fluorometer (Mithras LB 940; Berthold). The reaction was repeated after the addition of oligomycin (2 μ mol/L) as a mitochondrial ATPase inhibitor and of valinomycin (2.5 μ mol/L) as an oxidative phosphorylation inhibitor.²²

Ndufc2-silenced cells were washed once with PBS 1 \times and then incubated at 37°C in 1 mL of respiration buffer (as reported above for brain tissues) in the presence of both glutammate (2.5 mmol/L) and malate (2.5 mmol/L) or succinate (5 mmol/L) for 60 minutes. Cells were washed once with ice-cold PBS and then lysed with 90 μ L of ATP-lysis buffer (Tris HCl [pH 7.5] 200 mmol/L, NaCl 2 mol/L, EDTA [pH 7.5] 20 mmol/L, and Triton X-100 0.2%). Lysates were collected on ice and centrifuged at 4°C at 12 000 rpm for 15 minutes. Ten microliters of supernatant from each sample was plated in a dark 96-multiwell plate; 100 μ L of total ATP-kit buffer from the kit were added to each well. Chemiluminescent signal was read with the fluorometer. Five microliters of cell lysate were used for protein concentration and ATP-level normalization. The reaction was repeated after the addition of oligomycin (2 μ mol/L) and valinomycin (2.5 μ mol/L), as described above. Experiments were performed in triplicate.

Table 1. Modulation of Inflammatory Genes in *Ndufc2*-Silenced A10 Cells

Gene Name	Control	RNAi1	RNAi2
Ccl2	−0.83 \pm 0.77	1.05 \pm 0.83	0.73 \pm 0.98
Ccl6	−6.70 \pm 15.4	1.80 \pm 11.1	10.61 \pm 2.6
Ccr5*	86.30 \pm 17*	−42.24 \pm 21*	−8.31 \pm 15*
Ccr6*	62.44 \pm 1.3*	−1.40 \pm 2.5*	−1.23 \pm 1.9*
Cxcl10	0.86 \pm 3.4	−8.26 \pm 6.5	−3.25 \pm 5
Il10	−0.78 \pm 20	0.63 \pm 14	13.86 \pm 3.9
Il10ra	3.46 \pm 21	9.49 \pm 18	16.40 \pm 2.5
Il13*	1.81 \pm 51*	8.27 \pm 38*	36.22 \pm 8.5*
Il13ra1*	3.67 \pm 22*	42.77 \pm 26*	22.81 \pm 2*
Il15*	−28.93 \pm 1.9*	3.04 \pm 2.1*	1.89 \pm 0.15*
Lta	3.17 \pm 74	37.17 \pm 64	58.72 \pm 2.1
Tnf*	2.58 \pm 12.9*	9.37 \pm 11.9*	10.81 \pm 1.0*
Tgfb1*	−531 \pm 3.85*	3.57 \pm 3.8*	3.38 \pm 0.25*
Tnfrsf1b*	−11.04 \pm 3*	3.73 \pm 3.1*	2.77 \pm 0.18*

Ccl2 indicates chemokine ligand 2; Ccl6, chemokine ligand 6; Ccr5, chemokine receptor type 5; Ccr6, chemokine receptor type 6; Cxcl10, chemokine ligand 10; Il10, interleukin-10; Il10ra, interleukin-10 receptor alpha; Il13, interleukin-13; Il13ra1, interleukin-13 receptor alpha 1; Il15, interleukin-15; Lta, lymphotoxin-alpha; Tgfb1, transforming growth factor beta 1; Tnf, tumor necrosis factor; Tnfrsf1b, tumor necrosis factor receptor superfamily, member 1B.

*Genes markedly modulated by *Ndufc2* silencing (number of experiments=3).

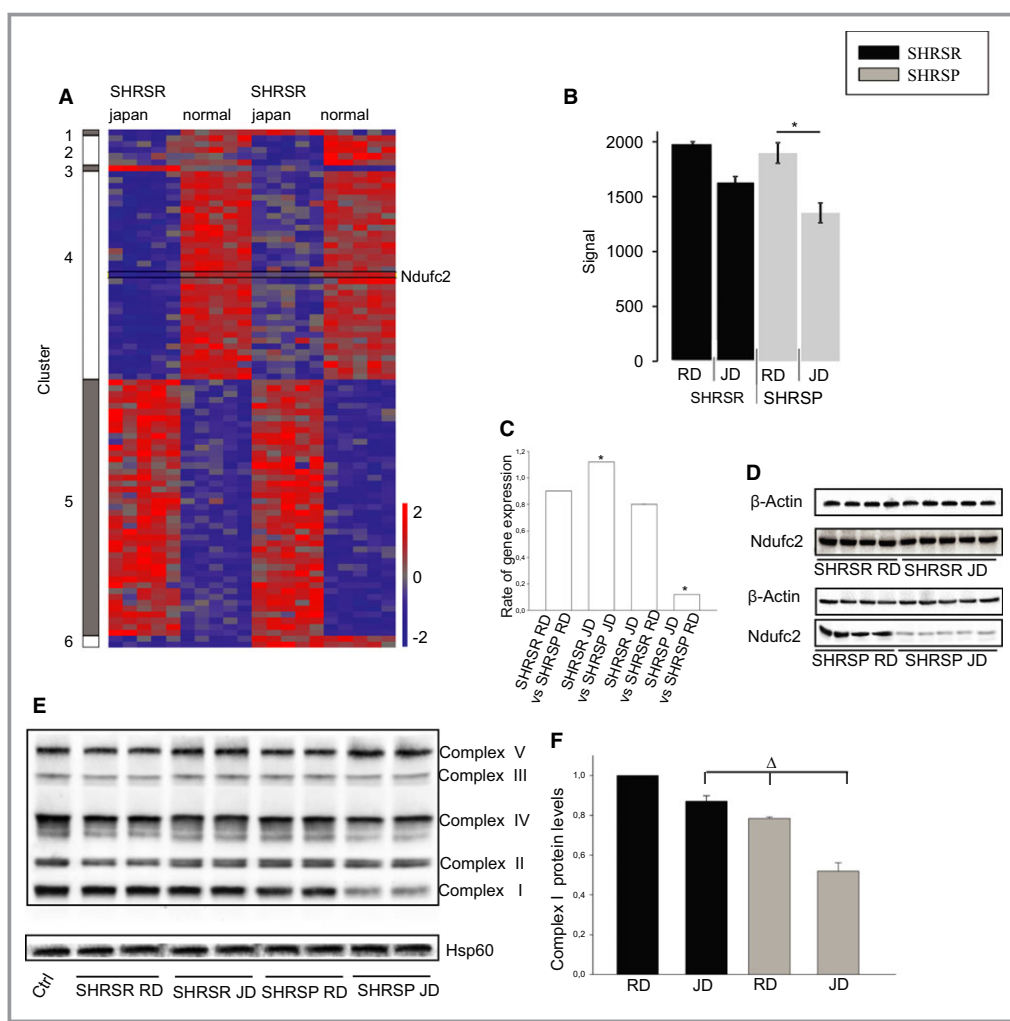


Figure 1. *Ndufc2* is differentially modulated in brains of SHRSP under stroke-permissive diet. Evidence of complex I dysfunction in brains of SHRSP. A, Heatmap represents expression differences between samples in a color scale ranging from red (up-regulated) to blue (down-regulated). Probe sets representing *Ndufc2* are marked. Two-way ANOVA analysis revealed significant *Ndufc2* effects in strain ($P=6.96E^{-05}$), diet ($P=8.29E^{-10}$), and interaction ($P=2.79E^{-03}$; $n=5$ for each strain at each diet). Corresponding densitometric analysis for *Ndufc2* expression levels in brains of SHRSR and SHRSP under either RD or JD is shown in (B). C, Confirmation of *Ndufc2* differential expression by RT-PCR in the same tissue samples; $*P<0.0001$ for each ratio indicated in the figure ($n=6$ for each group). D, Representative western blot of *Ndufc2* expression levels, confirming reduced expression in brains of SHRSP under JD. E, Representative western blot of BNG ($n=3$) and corresponding densitometric analysis (F). $\Delta P<0.01$ for SHRSP JD versus both SHRSR RD and SHRSR JD. G, Complex I activity measurement; (H) mitochondrial membrane potential assessment. $*P<0.0001$ for SHRSP JD versus all other samples (I). ATP-level measurement in the same tissue samples. Oligomycin was used to confirm functionality of ATP assay as an inhibitor of oxidative phosphorylation. Valinomycin was used as a mitochondrial ATPase inhibitor. $*P<0.0001$ for SHRSP JD versus all other samples and for each inhibited experimental point versus the noninhibited corresponding sample ($n=6$ for each group). Values are expressed as means \pm SD. Ctrl indicates control; Hsp60, heat shock protein 60; JD, Japanese-style stroke-permissive diet; RD, regular diet; RT-PCR, reverse-transcriptase polymerase chain reaction; SHRSP, stroke-prone spontaneously hypertensive rat; SHRSR, stroke-resistant SHR.

Generation of a Genetically Manipulated Animal Model

Generation of SHR-*Ndufc2* mutant strains (SHR-*Ndufc2*^{em1M_{cwi}} and SHR-*Ndufc2*^{em2M_{cwi}}) was performed at the Medical

College of Wisconsin (Milwaukee, WI) under protocols approved by the institutional animal care and use committee. Generation of ZFN mutants was performed as previously described.^{23,24} ZFN constructs specific for rat *Ndufc2* were designed, assembled, and validated by Sigma-Aldrich to the

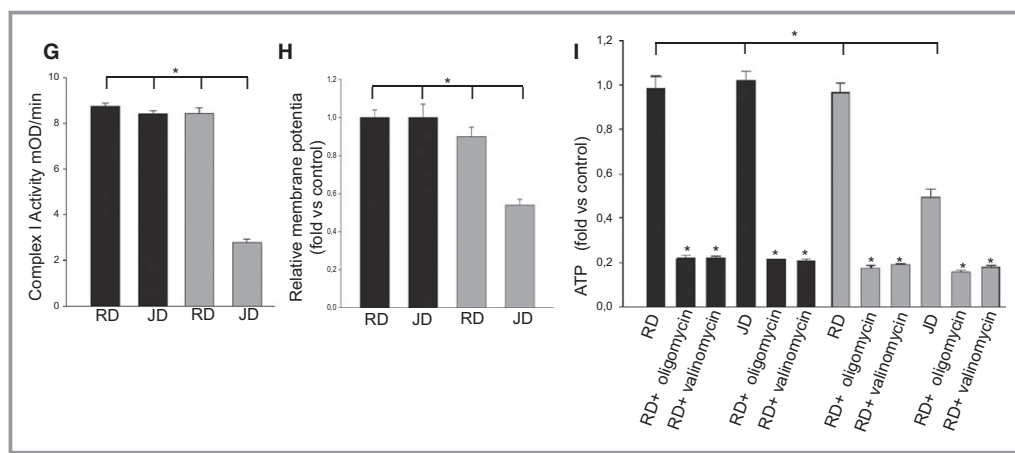


Figure 1. Continued.

exon 1 sequence GGCTTCCTGGGCTACTGCacgggcCTGATGGA-CAACATG where each ZFN monomer binds the underlined sequence on opposite strands. In vitro-transcribed ZFN mRNA SHR/NCrl (SHR) rat embryos were transferred to pseudopregnant Sprague-Dawley female rats to be carried to parturition. The resulting pups were ear punched and DNA was extracted and screened for ZFN-induced mutations by the Surveyor Nuclease assay (Transgenomic, Inc., Omaha, NE), as described previously,²³ using primers flanking the target sequence (*Ndufc2*_F: 5'-CGCATCAATATGATGAACGG-3' and *Ndufc2*_R: 5'-CGCTGAAACTCTAGACGGG-3'). Two positive mutant founders were identified and confirmed to have ZFN-induced mutations. Sanger sequencing confirmed mutant alleles within exon 1 in each founder, a 9-bp deletion (del-9 ACGGGCCTG; *Ndufc2*-m1) and a net 107-bp deletion (del-111 GCCCGTGCACTAGCCCAGGAAGCCGATGTAGACAAGCCGCGG GTCGTTTCAGCTTGGGCGGGGGCAGTCTCCGGGCCTCATCCGGC AAGAATCTTAAGGGCTCATGGCCCGG plus ins-4 TTGT; *Ndufc2*-m2). The 2 founders were back-crossed to the SHR/NCrl strain to establish the SHR-*Ndufc2*^{em1M_{cwi}} and SHR-*Ndufc2*^{em2M_{cwi}} breeding colonies, respectively.

Both newborn rats and embryos obtained from colonies breeding were genotyped to assess allele carrier status for the corresponding deletion. Embryos were obtained at 7 to 8 days of gestation (SHR-*Ndufc2*^{em2M_{cwi}}, n=47 from 6 pregnant rats; SHR-*Ndufc2*^{em1M_{cwi}}, n=20 from 2 pregnant rats), and at 10 to 12 days of gestation (SHR-*Ndufc2*^{em2M_{cwi}}, n=8 from 2 pregnant rats; SHR-*Ndufc2*^{em1M_{cwi}}, n=8 from 2 pregnant rats).

For genotyping of SHR-*Ndufc2* mutant strains, genomic DNA was prepared from tail clipping of each animal by salt precipitation⁵ and subsequently used for PCR reactions.

For detection of the large 107-bp deletion (SHR-*Ndufc2*^{em2M_{cwi}}), PCR reactions were performed with 50 ng of genomic DNA in a total volume of 20 μ L containing

250 nmol/L of dNTP, 10 nmol/L of each primer (forward 5'-TCTTTGCCCCACTGTGGGTAT 3' and reverse 5'-CCCCTCCCCAGACCTTATGA-3'), and 0.3 U of Taq Polymerase (Life Technologies, Carlsbad, CA). They were processed on a thermal cycler system (T100 Thermal Cycler; Bio-Rad, Hercules, CA) by using "touchdown PCR" conditions (to increase specificity and reduce background amplification). Briefly, PCR was carried out at 95°C for 3 minutes followed by 14 cycles of denaturation at 95°C for 15 seconds, annealing at 63.5°C for 30 seconds with subsequent decreases of 0.5°C per cycle, and elongation at 72°C for 70 seconds, with subsequent 19 cycles at 95°C for 15 seconds, 56°C for 30 seconds, 72°C for 70 seconds, and a final extension of 72°C for 5 minutes. The resultant PCR product was a fragment of 318 bp in the wild-type strain and a fragment of 211 bp in the SHR-*Ndufc2*^{em2M_{cwi}} strain. They were resolved on 2.5% agarose gels and visualized using the ChemiDoc XRS+Imaging System (Bio-Rad).

For detection of the minor 9-bp deletion (SHR-*Ndufc2*^{em1M_{cwi}}), the PCR/single-strand conformational polymorphism analysis was used. Briefly, genotyping was carried out using 50 ng of DNA and the same set of oligonucleotides mentioned above. In this case, the forward primer was previously labeled with ATP- γ -³²P (PerkinElmer) by using T4 polynucleotide kinase (New England Biolabs, Boston, MA). PCR reactions were carried out with the same protocol described above, and products were loaded onto a 7% polyacrylamide gel run on a Gel Electrophoresis System (Thermo Fisher Scientific, Waltham, MA) for 4 hours. Images were acquired using the PMI Personal Molecular Imager System (Bio-Rad).

Because only heterozygous rats were obtained for both mutant lines, the molecular analyses and phenotypic studies were performed in heterozygous, compared to wild-type, rats of both SHR-*Ndufc2*^{em1M_{cwi}} and SHR-*Ndufc2*^{em2M_{cwi}} lines.

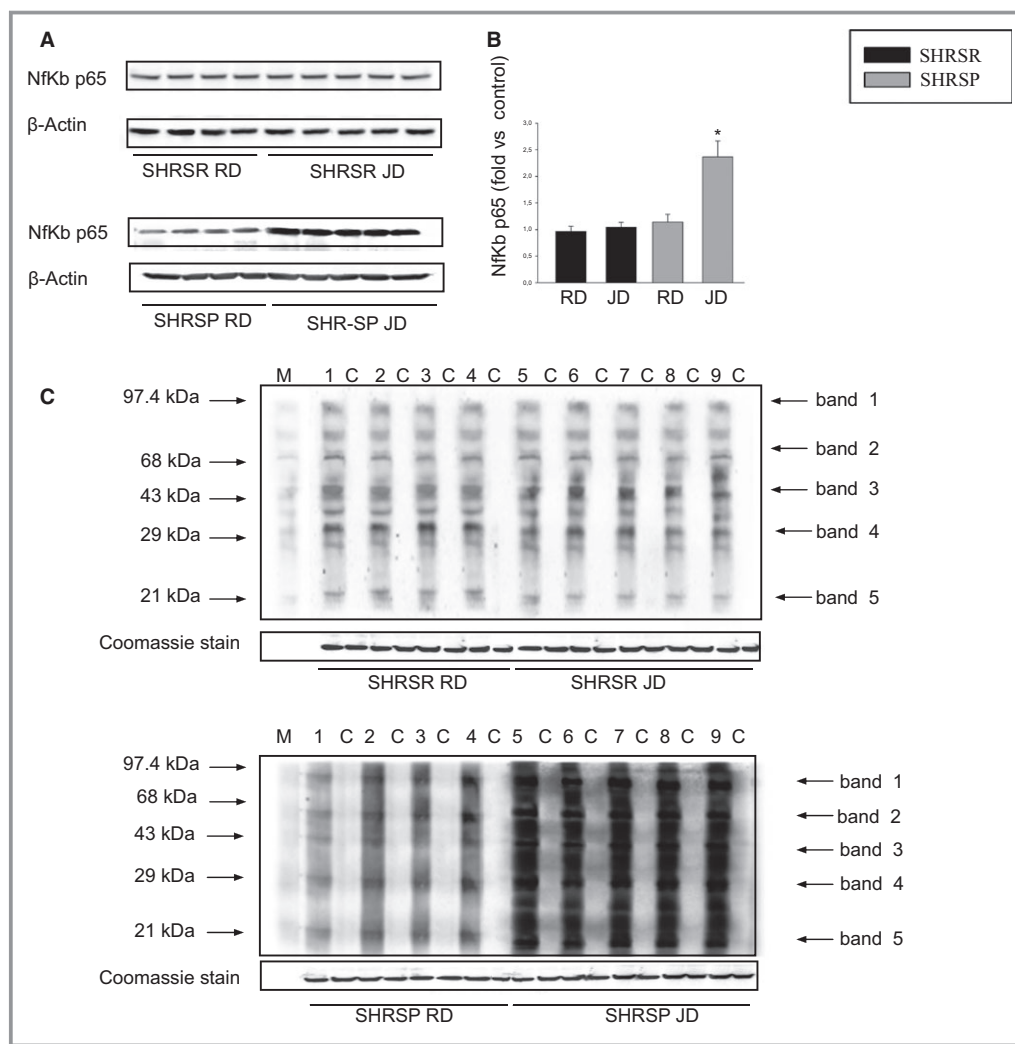


Figure 2. Molecular analyses in brains of SHRSP as compared to SHRSR under stroke-permissive diet. A, Representative western blot (n=3) of Nf-κBp65 in SHRSR and SHRSP upon either RD or JD with corresponding densitometric analysis (B). * $P<0.0001$ for SHRSP JD versus all other samples. D, Western blot of intracellular protein extracts immunostained for carbonylated proteins using the Oxyblot Protein Oxidation Detection kit (Millipore, Milan, Italy) in brains of SHRSR and SHRSP under either RD or JD. Each lane was loaded with 50 μ g of total proteins. Lane M, DNP marker. Each sample is run with its own untreated control (C). Normalization for lane protein loading was performed using Coomassie staining. D, Densitometric analysis of Oxiblot gels. Bar graphs represent chemiluminescence intensity relative to the gel loading band. Bands 1 to 5 refer to the most prominent bands on the blots (identified by arrows), whereas total refers to the total chemiluminescence intensity from all bands. * $P<0.0001$ for SHRSP JD versus SHRSR RD (total band). Densitometric values are expressed as means \pm SD. JD indicates Japanese-style stroke-permissive diet; Nf-κB, nuclear factor kappa B; RD, regular diet; SHRSP, stroke-prone spontaneously hypertensive rat; SHRSR, stroke-resistant SHR.

Phenotyping of SHR-*Ndufc2*^{em1M_{cwi}} and SHR-*Ndufc2*^{em2M_{cwi}} Lines

Feeding with JD was started at 6 weeks of age in both heterozygous and wild-type SHR-*Ndufc2*^{em1M_{cwi}} and SHR-*Ndufc2*^{em2M_{cwi}} rats, along with parental rats (for number of animals used, see Table 1). A group of rats (n=6) was sacrificed both at 6 weeks of age and at the end of 4 weeks

of either RD or JD for analysis of: *Ndufc2* by western blot and RT-PCR, SOD2, and Nf-κBp65 by western blot, complex I assembly and activity, mitochondrial membrane potential, ATP levels, and tissue oxidative stress. Remaining rats were subsequently monitored up to 3 months for signs of renal damage (detected by 24-hour proteinuria measurement in metabolic cages) and of stroke (by evidence of paresis, paralysis, hemiplegia, epilepsy, or sudden death). Animal

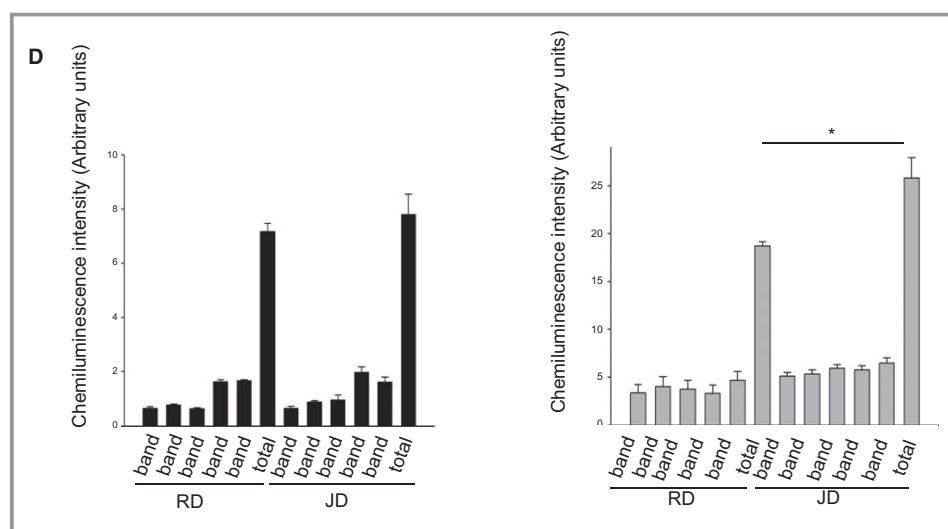


Figure 2. Continued

studies were approved by the Neuromed Institutional review committee. Procedures involving animals and their care were carried out in accord with our institution's guidelines, which comply with national and international laws and policies.

Genotyping of NDUFC2 Markers in an Italian Cohort of Early-Onset Ischemic Strokes and Controls

Study population

Four hundred eighty-four unrelated Caucasian young adults (<65 years) who were referred to the (1) Angelo Bianchi Bonomi Hemophilia and Thrombosis Center of the University of Milan and (2) to Department of Neurology, Sapienza University, Sant'Andrea Hospital, Rome, for thrombophilia screening after a first ischemic stroke were enrolled in this study. Patient and control populations were in part or in whole previously investigated in relation to polymorphisms of genes encoding procoagulant and inflammatory factors and of methionine metabolism components, chromosome 12p13, and NPR3 gene promoter polymorphisms.^{25–28} Median time interval between ischemic stroke and blood sampling was 6 months (range, 1 month–10 years); 75% of patients had blood sampled within 2 years and 64% within 1 year after the event. Clinical records were reviewed, and when the type of stroke was not specified, neurologists who took care of patients during the acute phase were contacted. Clinical diagnosis was objectively confirmed by computed tomography scans or magnetic resonance or magnetic resonance angiography and intra-arterial angiography. According to Trial of ORG 10172 in Acute Stroke Treatment (TOAST) classification, 14.7% of enrolled patients were classified as large artery stroke, 5.2% as cardioembolic,

16.8% as small artery (lacunar), 51.8% as undetermined etiology, and 11.6% as other determined etiology stroke. All patients underwent a cardiologic evaluation, transthoracic echocardiography, and Doppler examination of neck vessels.

One thousand one hundred sixty-five healthy unrelated Caucasian subjects comparable for age and sex were chosen from the whole population of controls made of partners and friends who accompanied patients to the Center in the same period as patients and agreed to be investigated. Previous thrombosis in the controls was excluded using a validated structured questionnaire.²⁹ The presence, at the time of stroke for patients and at the time of blood sampling for controls, of hypertension, hypercholesterolemia, diabetes mellitus, and smoking habit (at least 5 cigarettes daily) was recorded.

The study was approved by the institutional review board of the University of Milan, University of Rome, and University of Florence, and all subjects gave their informed consent to the study.

Genomic DNA extraction

Genomic DNA was isolated from venous peripheral blood by using the FlexiGene kit (Qiagen). Quality and concentration of DNA were determined by a NanoDrop1000 spectrophotometer (Thermo Fisher Scientific).

Genotyping

We studied 3 tagSNPs in NDUFC2: rs641836, rs584981, and rs11237379 by 7900HT real-time PCR and TaqMan technology assays (c__26529993_10, c__614632_10 and c__2999825_10, respectively; Life Technologies). The 3 NDUFC2 tagSNPs were selected for analysis by using the algorithm-Tagger-pairwise Tagging (HapMap database and software, <http://>

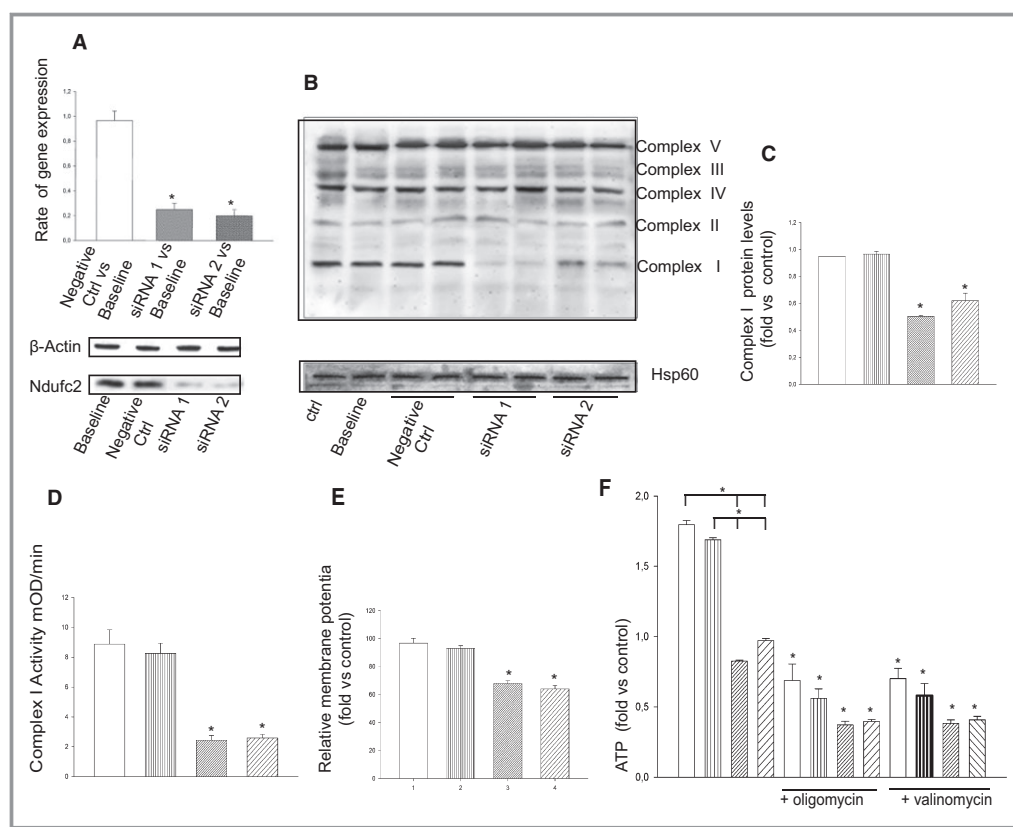


Figure 3. In vitro *Ndufc2* silencing in A10 cells. A, Confirmation of *Ndufc2* silencing by RT-PCR and western blot; $*P<0.0001$ for each siRNA versus baseline ($n=6$). B, Representative western blot of BNG ($n=3$) in silenced versus nonsilenced A10 cells. All complexes belonging to OXPHOS are identified on the right side; Hsp60 was used to normalize protein expression level. C, Densitometric analysis of BNG. $*P<0.0001$ for each siRNA versus both baseline and negative CTRL. D, Assessment of complex I activity. E, Mitochondrial membrane potential assessment. $*P<0.0001$ for each siRNA versus both baseline and negative CTRL ($n=6$). F, Measurement of ATP levels in silenced versus nonsilenced cells and both in the absence and in the presence of oligomycin and valinomycin (to confirm functionality of the ATP assay as specified above). $*P<0.0001$ for each siRNA versus baseline and versus negative CTRL, and for each inhibited point versus the noninhibited corresponding sample ($n=6$ for each group). G, Assessment of reactive oxygen species levels; $*P<0.0001$ for each siRNA versus both baseline and negative CTRL ($n=6$ for each group). H, Representative western blot ($n=3$) of $\text{Nf-}\kappa\text{Bp65}$ and of its inhibitor, $\text{I}\kappa\text{B}$, in silenced versus nonsilenced cells, showing that increases of $\text{Nf-}\kappa\text{Bp65}$ expression are accompanied by decreases of $\text{I}\kappa\text{B}$ expression levels. I, FACS analysis and corresponding percent values of necrotic cells (J). $*P<0.0001$ for each siRNA versus both baseline and negative CTRL ($n=5$). Values are expressed as means \pm SD. Ctrl indicates control; DCHF, 2',7'-dichlorodihydrofluorescein; FACS, fluorescence-activated cell sorting; Hsp60, heat shock protein 60; $\text{I}\kappa\text{B}$, nuclear factor of kappa light polypeptide gene enhancer in B-cells inhibitor; $\text{Nf-}\kappa\text{B}$, nuclear factor kappa B; RT-PCR, reverse-transcriptase polymerase chain reaction.

hapmap.ncbi.nlm.nih.gov) on *NDUFC2* gene for CEU population. They captured 100% of alleles with a mean r^2 of 0.961. Polymorphisms information were assessed in the Single Nucleotide Polymorphism (dbSNP) NCBI (<http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?db=snp&cmd=search&term=>), and ENSEMBL (<http://www.ensembl.org/index.html>) databases.

To assess functional relevance of the rs11237379 variant allele, we characterized *NDUFC2* expression by RT-PCR in a cohort of young healthy subjects (mean age, 35 ± 6 ; females,

52%), free from cardiovascular risk factors, carrying wild-type ($n=6$), heterozygous ($n=12$), and mutant homozygous ($n=6$) genotypes. Lymphocyte extraction was performed by following standard procedures with Ficoll.³⁰ RNA extraction and cDNA were obtained as described above. Finally, RT-PCR of human *NDUFC2* was performed with the following set of oligonucleotides: forward 5'-CCTGTATGCTGTGAGGGACC-3' and reverse 5'-CGACGTTTCAGCTCCACAACA-3'. β -actin: forward 5'-GCAAGAGATGGCCACGGCTG-3' and reverse 5'-CCACAGGACTCCATGCCAG-3'.

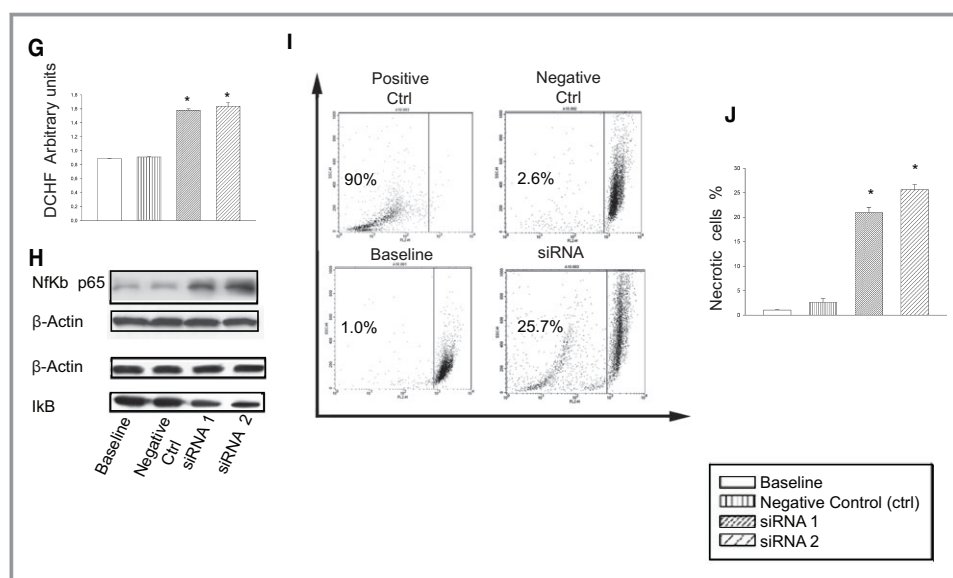


Figure 3. Continued

Statistical Analysis

In vitro and in vivo experiments

Data of *Ndufc2* expression levels, western blot densitometric analyses, complex I activity and ATP levels, results of FACS, rat body weight, systolic blood pressure, and urinary protein levels are provided as means±SD. Comparisons between 2 groups were performed by using *t* test analyses. When more than 2 groups were compared at the same time, 1-way ANOVA followed by the Bonferroni post-hoc test was performed. Survivor function in rats monitored over JD feeding was estimated by the life-table method. Log-rank and Wilcoxon statistics were used for testing equality of survivor functions.

SPSS statistical software (version 12.0; SPSS, Inc., Chicago, IL) was used for statistical analysis. Statistical significance was stated at the $P<0.05$ level.

Human genetic analyses

Statistical analysis was performed using the SPSS package (v20). Hardy–Weinberg equilibrium (HWE) was evaluated by the chi-square (χ^2) test. Genotype distributions were compared between patients and controls or between TOAST types by χ^2 analysis. Categorical variables are expressed as frequencies and percentages. Unless otherwise indicated, continuous data are given as median and interquartile range (IQR). Comparisons of continuous variables between patients and controls or among genotypes were performed by the nonparametric Mann–Whitney or Kruskal–Wallis test.

Multiple logistic regression analysis was used to estimate odds ratios (ORs) and 95% CIs for risk of stroke under the assumption of an additive, as well as a dominant and a recessive effect of each allele. To evaluate whether polymor-

phisms were independently associated with stroke, multiple logistic regression analyses were adjusted for traditional cardiovascular risk factors (age, sex, hypertension, diabetes mellitus, dyslipidemia, and smoking habit). A value of $P<0.05$ was chosen as the cut-off level for statistical significance.

Results

Identification of *Ndufc2* as a Sequence Down-regulated in Brains of SHRSP Under JD and Related Molecular Analyses

Eight-six probe sets were revealed significantly differentially modulated inside the STR1/QTL. After applying a K-mean clustering with $K=6$ and the 2-way ANOVA statistical analysis, as specified in the Methods section, we detected the gene encoding *Ndufc2* (NADH dehydrogenase [ubiquinone] 1, subcomplex unknown, 2), mapping 8 Mb apart from the Lod score peak of the QTL, as the sequence differentially modulated, significantly down-regulated, in the brain of SHRSP versus SHRSR when fed with JD (P values for significant *Ndufc2* effects in strain [$P=6.96E^{-05}$], diet [$P=8.29E^{-10}$], and interaction [$P=2.79E^{-03}$]; $n=5$ for each strain at each diet; Figure 1A and 1B).

Analysis of *Ndufc2* mRNA expression and protein levels confirmed the microarray differential expression results ($P<0.0001$; $n=6$; Figure 1C and 1D).

Ndufc2 is a subunit of OXPHOS complex I. Therefore, we evaluated mitochondrial function in brains of SHRSP and SHRSR under JD. Brains of JD-fed SHRSP showed significant reduction of both complex I assembly ($P<0.01$; $n=3$) and activity ($P<0.0001$; $n=6$; Figure 1E through 1G), mitochondrial membrane potential ($P<0.0001$; $n=6$; Figure 1H), and ATP

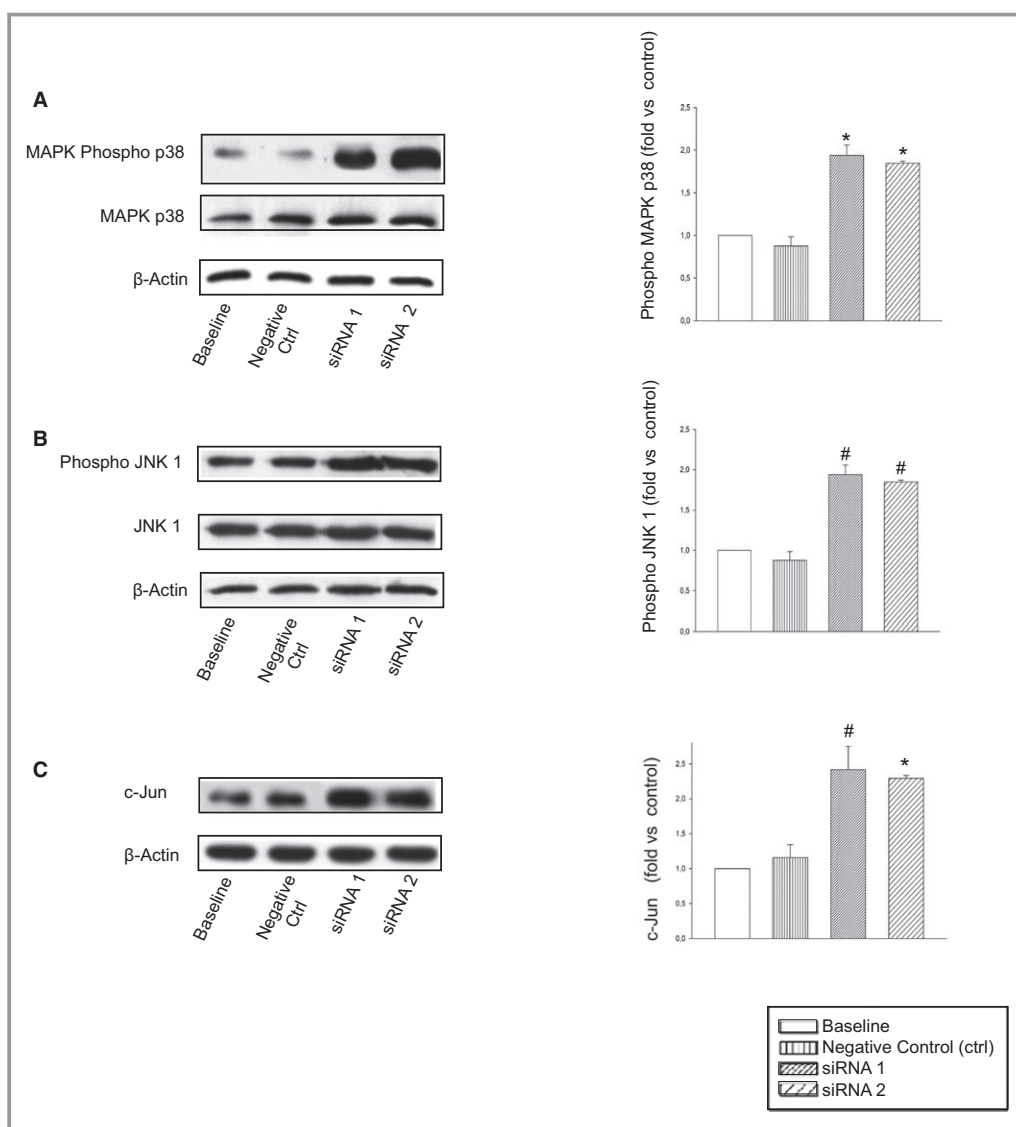


Figure 4. Expression levels of proteins involved in necrosis in *Ndufc2*-silenced as compared to non-silenced A10 cells. A, MAPKp38; (B) JNK1; and (C) c-Jun. Number of western blots for each protein=3. Densitometric analyses are reported on the right side. * $P<0.0001$ and # $P<0.001$ for each siRNA versus baseline and negative CTRL. Ctrl indicates control; JNK1, c-Jun N-terminal kinase 1; MAPK, mitogen-activated protein kinase.

production ($P<0.0001$; $n=6$; Figure 1I), as well as remarkable increases of both inflammation ($P<0.0001$; $n=3$) and oxidative stress ($P<0.0001$; $n=3$; Figure 2), likely as a consequence of mitochondrial uncoupling.

Next-generation sequencing, covering full exons, introns, and 10 kb upstream and downstream on either side of *Ndufc2*, did not show variations between SHRSP and SHRSR.

In Vitro Effects of *Ndufc2* Silencing

We then tested whether *Ndufc2* knockdown in vascular cells could recapitulate the cellular abnormalities observed in

SHRSP under JD. *Ndufc2* silencing was confirmed by both RT-PCR and western blotting ($P<0.0001$; $n=6$; Figure 3A). In this experimental condition, complex I assembly ($P<0.0001$; $n=3$) and activity ($P<0.0001$; $n=6$), mitochondrial membrane potential ($P<0.0001$; $n=6$), and ATP production ($P<0.0001$; $n=6$) were significantly decreased (Figure 3B through 3F). Reactive oxygen species (ROS) production ($P<0.0001$; $n=6$), Nf- κ B signaling, and other markers of inflammation increased in *Ndufc2*-silenced cells (Figure 3G and 3H; Table 1). Fluorescence-activated cell sorting (FACS) analysis revealed a significant decrease of cell viability and a significant increase of cell necrosis ($P<0.0001$; $n=5$; Figure 3I and 3J). Expression

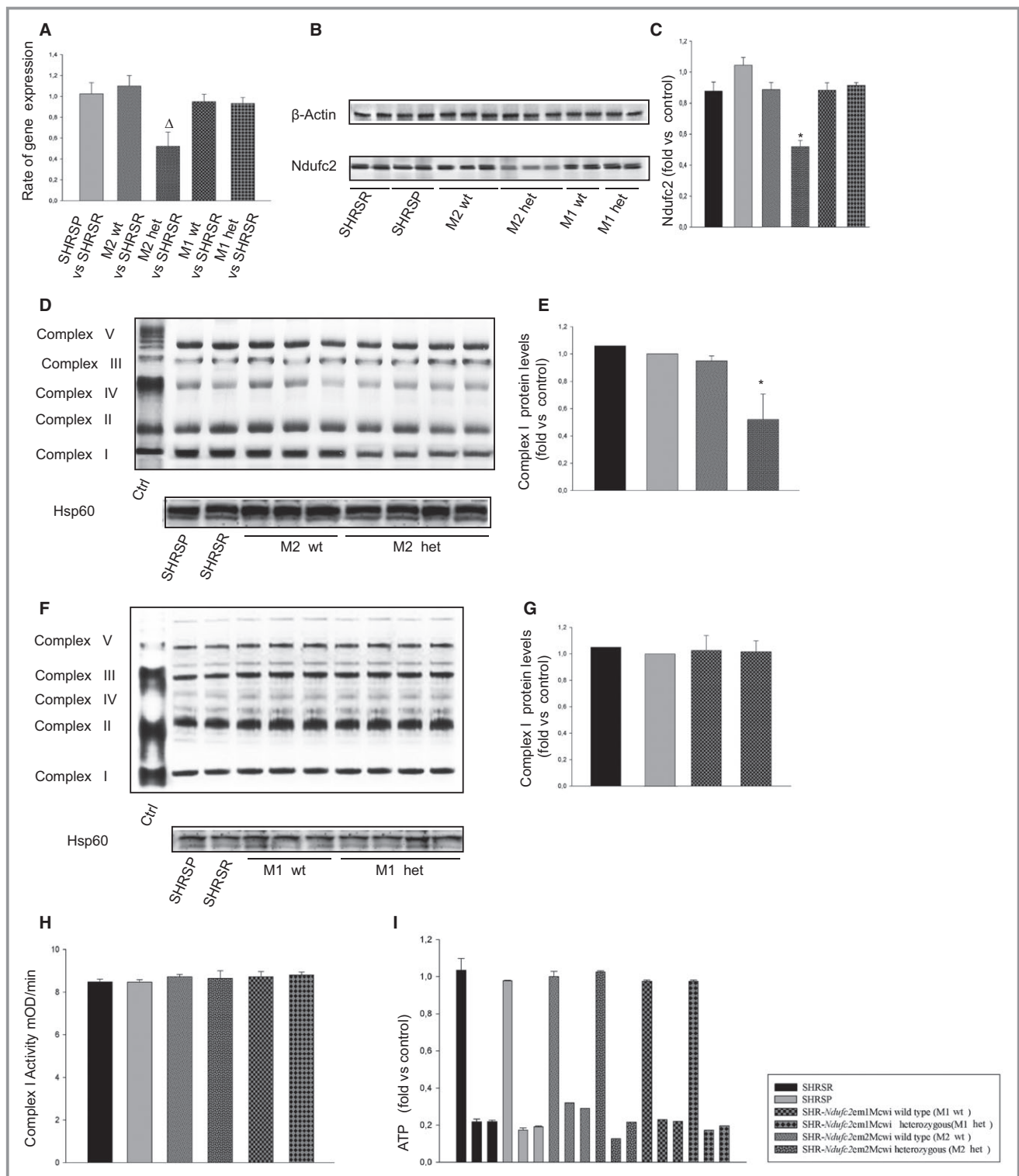


Figure 5. Molecular analyses of brains of all strains at 6 weeks of age. A through C, Brains of heterozygous SHR-*Ndufc2*^{em2M_{cwi}} showed significantly lower amount of *Ndufc2* mRNA and protein expression, as compared to both parental SHRSP and SHR-*Ndufc2*^{em1M_{cwi}}. ^Δ*P*<0.01 versus SHRSP; **P*<0.0001 versus all other samples (*n*=6 for each group). D through G, Representative western blot of BNG (*n*=3) showing defective complex I assembly only in heterozygous SHR-*Ndufc2*^{em2M_{cwi}} rats. Corresponding densitometric analyses are shown on the right side of each blot. **P*<0.0001 versus all other samples. H, Complex I activity and (I) ATP levels were comparable among all strains at this age (*n*=3). Values are expressed as means±SD. Ctrl indicates control; het, heterozygous; Hsp60, heat shock protein 60; SHRSP, stroke-prone spontaneously hypertensive rat; SHRSP, stroke-resistant SHR; wt, wild type.

Table 2. SBP, BW, and Proteinuria Levels in Parental and Genetically Manipulated Rat Lines Upon JD Feeding

	BW (g)			SBP (mm Hg)			Proteinuria (mg/24 h)					
	4 Weeks	7 Weeks	8 Weeks	12 Weeks	4 Weeks	7 Weeks	8 Weeks	12 Weeks	4 Weeks	7 Weeks	8 Weeks	12 Weeks
SHRSR	220±13 (n=18)	—	260±16 (n=18)	289±8* (n=18)	180±4 (n=18)	—	189±6 (n=18)	188±4 (n=18)	33±5.5 (n=18)	—	68±14 (n=18)	85±16* (n=18)
SHRSP	177±6† (n=18)	192±6‡ (n=5)	—	—	181±14 (n=18)	193±14 (n=5)	—	—	188±12† (n=18)	240±30‡ (n=5)	—	—
SHR <i>Ndufc2</i> ^{pm1Mowi} w.t.	235±15 (n=10)	—	287±12 (n=10)	309±27* (n=10)	177±7 (n=10)	—	183±2 (n=10)	188±2 (n=10)	27±12 (n=10)	—	56±16 (n=10)	67±6* (n=10)
SHR <i>Ndufc2</i> ^{pm1Mowi} hetero	235±14 (n=17)	—	281±18 (n=17)	308±21* (n=17)	174±6 (n=17)	—	178±6 (n=17)	182±7 (n=17)	24±7 (n=17)	—	43±6 (n=17)	71±19* (n=17)
SHR <i>Ndufc2</i> ^{pm2Mowi} w.t.	239±13 (n=12)	—	299±16 (n=12)	314±49* (n=12)	181±3 (n=12)	—	181±9 (n=12)	190±3 (n=12)	31±5 (n=12)	—	47±26 (n=12)	72±22 Δ (n=12)
SHR <i>Ndufc2</i> ^{pm2Mowi} hetero	237±27 (n=26)	—	293±30 (n=24)	312±36* (n=16)	177±7 (n=26)	—	189±2 (n=24)	186±6 (n=16)	84±35§ (n=26)	—	146±58 ¶ (n=24)	126±27 (n=16)

SHRSP survived up to 7 weeks of JD feeding. BW indicates body weight; JD, Japanese-style stroke-permissive diet; SBP, systolic blood pressure; SHRSP, stroke-prone spontaneously hypertensive rat; SHRSR, stroke-resistant SHR.

**P*<0.0001 for 12 weeks vs 4 weeks for each strain.

†*P*<0.001 vs other strains at 4 weeks.

‡*P*<0.0001 for SHRSP/7 weeks vs SHRSP/4 weeks.

§*P*<0.05 for heterozygous M2/4 weeks vs wild-type M2, heterozygous, and wild-type M1/4 weeks.

||*P*<0.0001 for heterozygous M2/8 weeks vs all other strains/8 weeks.

Δ*P*<0.05 for heterozygous M2/8 weeks vs heterozygous M2/4 weeks.

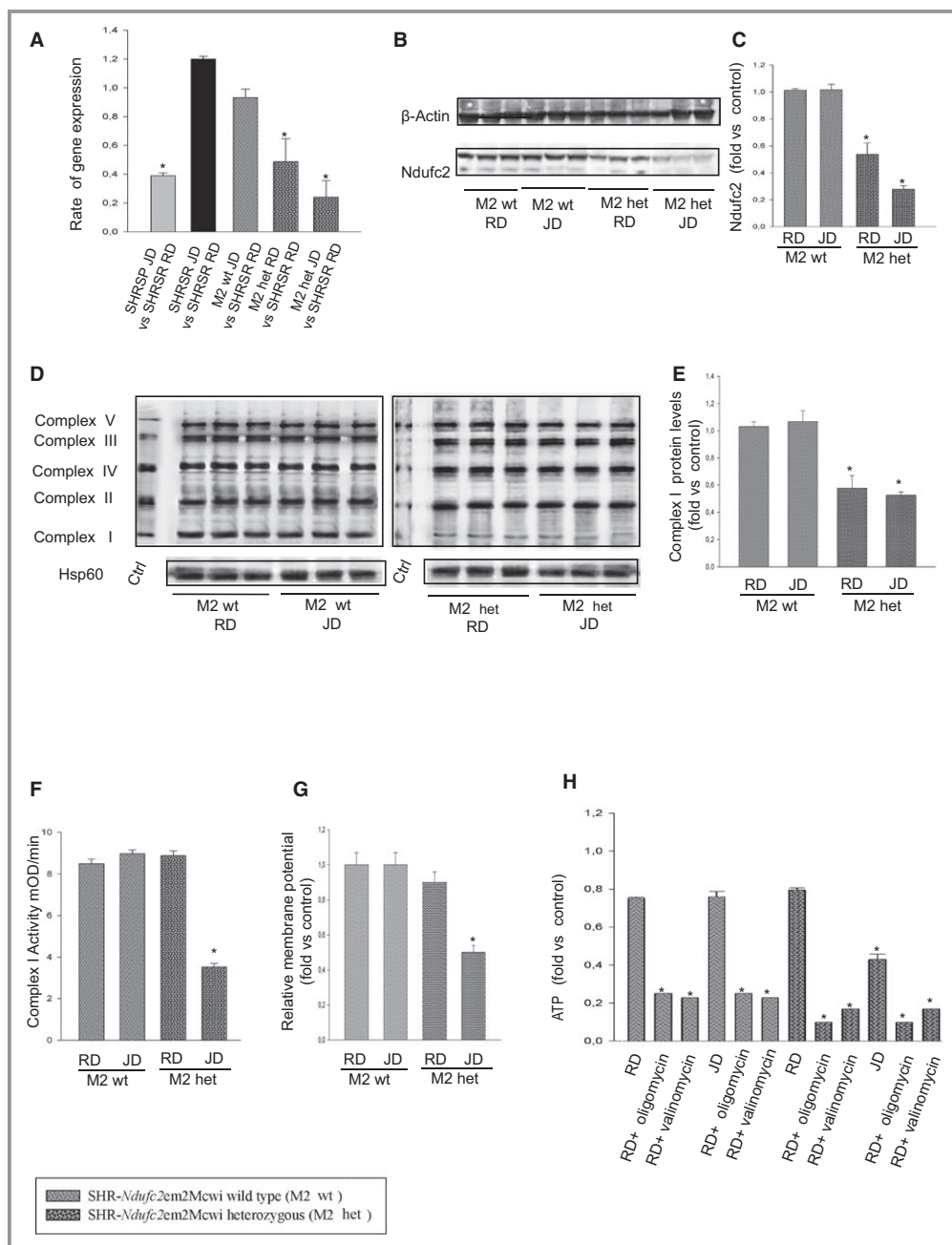


Figure 6. Molecular analyses in brains of both wild-type and heterozygous SHR-*Ndufc2*^{em2Mcowi} lines at the end of 4 weeks exposure to either RD or JD. A, Brain *Ndufc2* expression in both wild-type and heterozygous SHR-*Ndufc2*^{em2Mcowi} is compared to that of RD-fed parental SHRSR; **P*<0.0001 for each indicated ratio (*n*=5). B, *Ndufc2* protein expression levels in brains of wild-type and heterozygous SHR-*Ndufc2*^{em2Mcowi} (number of western blots=3) with corresponding densitometric analysis (C); **P*<0.0001 for heterozygous SHR-*Ndufc2*^{em2Mcowi} upon RD and JD versus wild type. D and E, Representative western blots of BNG in wild-type and heterozygous SHR-*Ndufc2*^{em2Mcowi} (*n*=3 each) with corresponding densitometric analysis. **P*<0.0001 for heterozygous SHR-*Ndufc2*^{em2Mcowi} upon RD and JD versus wild type. F, Complex I activity measurement; (G) mitochondrial membrane potential assessment; **P*<0.0001 for heterozygous SHR-*Ndufc2*^{em2Mcowi} upon JD versus RD and versus wild-type line at both RD and JD (*n*=5 each). H, ATP levels measurement in both wild-type and heterozygous rat lines upon RD or JD (*n*=5 each). **P*<0.0001 for heterozygous SHR-*Ndufc2*^{em2Mcowi} versus wild-type line and for each inhibited point versus the not inhibited corresponding sample. Values are expressed as means±SD. BNG indicates Blue Native-PAGE; Ctrl, control; het, heterozygous; Hsp60, heat shock protein 60; JD, Japanese-style stroke-permissive diet; RD, geular diet; SHRSP, stroke-prone spontaneously hypertensive rat; SHRSR, stroke-resistant SHR; wt, wild type.

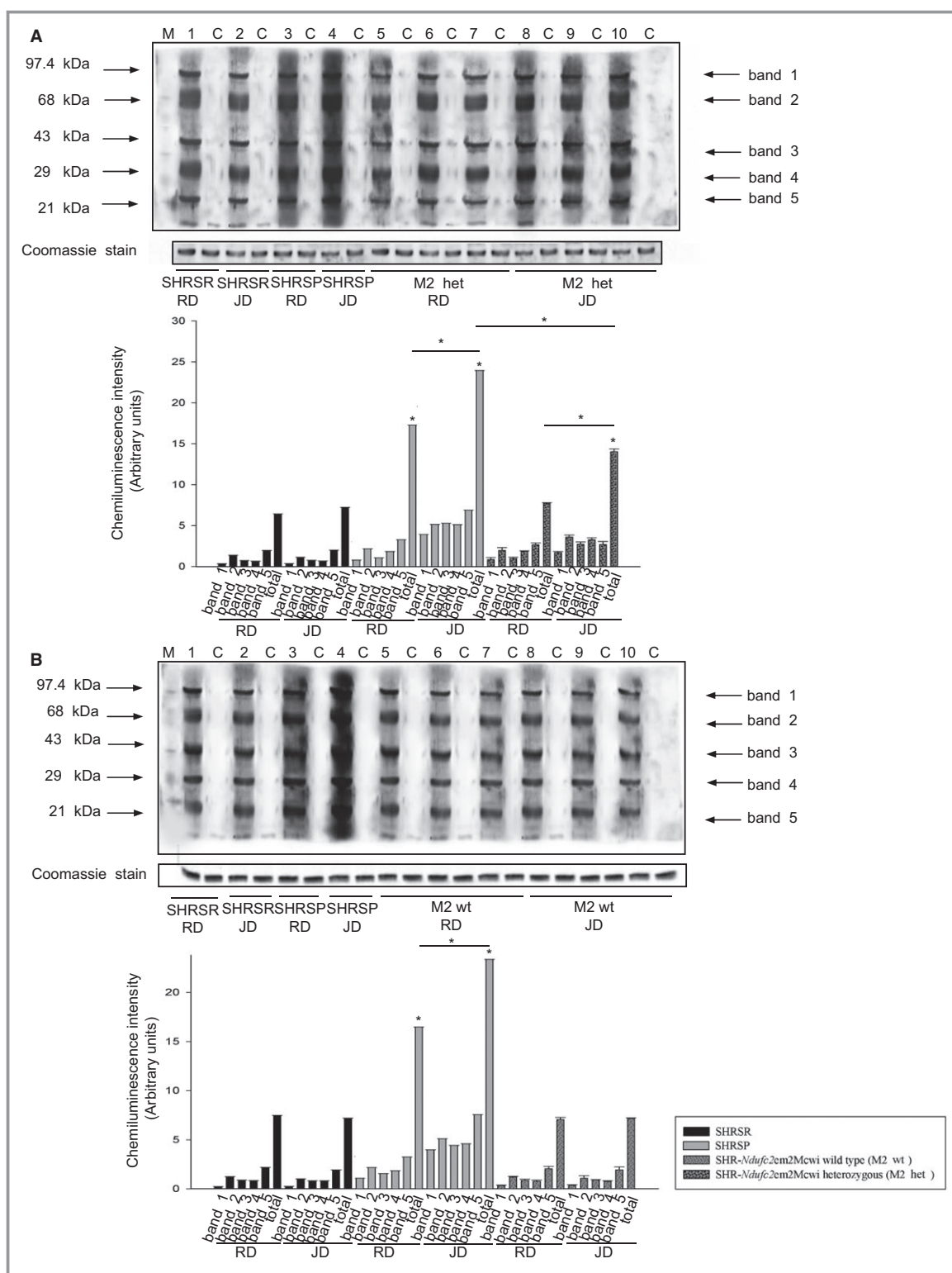


Figure 7. Detection of oxidative stress in brains of SHR-*Ndufc2*^{em2Mcowi} under either RD or JD. A, Detection of carbonylated proteins in brains of heterozygous SHR-*Ndufc2*^{em2Mcowi} at the end of 4 weeks of either RD or JD feeding with corresponding densitometric analysis (graph below). * $P < 0.0001$ for SHRSP versus RD and for heterozygous SHR-*Ndufc2*^{em2Mcowi} JD versus RD, as well as versus SHRSP JD. B, Detection of carbonylated proteins in brains of wild-type SHR-*Ndufc2*^{em2Mcowi} at the end of 4 weeks of either RD or JD feeding with corresponding densitometric analysis (graph below); * $P < 0.0001$ for SHRSP JD versus RD and for both versus all other strains (total band). Het indicates heterozygous; JD, Japanese-style stroke-permissive diet; RD, geular diet; SHRSP, stroke-prone spontaneously hypertensive rat; SHRSR, stroke-resistant SHR; wt, wild type.

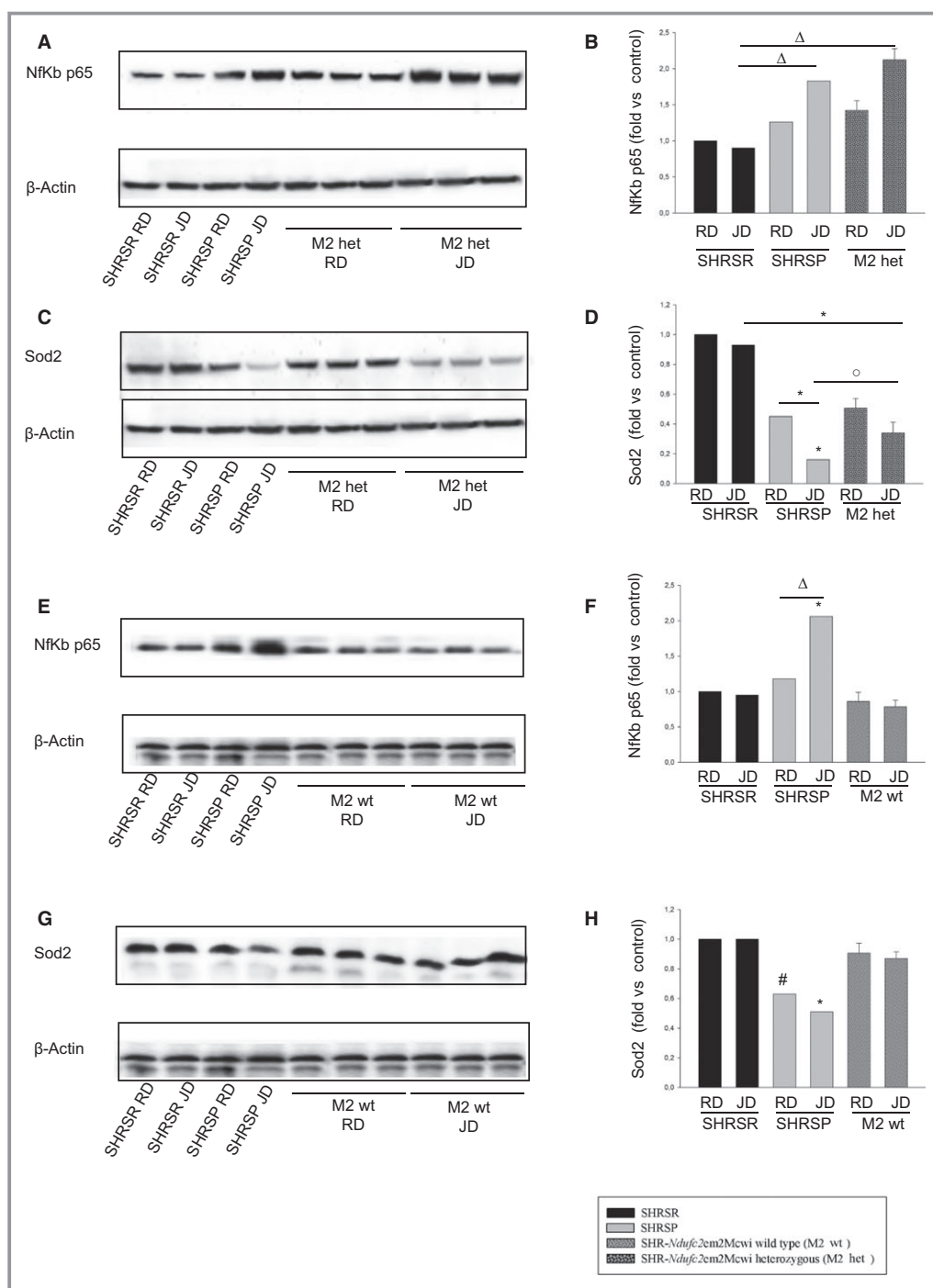


Figure 8. Molecular analyses in brains of wild-type and heterozygous SHR-*Ndufc2*^{em2Mcowi} at the end of either RD or JD exposure. A through D, Representative western blots of Nf-kBp65 and of SOD2 (n=3) with corresponding densitometric analysis in brains of heterozygous SHR-*Ndufc2*^{em2Mcowi}, as compared to parental, lines at the end of 4 weeks of either RD or JD feeding. E through H, Same analyses (n=3) were performed in brains of wild-type SHR-*Ndufc2*^{em2Mcowi}. Densitometric values are expressed as means±SD. Significance values: (B) $\Delta P<0.01$ for SHRSP JD and heterozygous SHR-*Ndufc2*^{em2Mcowi} JD versus SHRSR JD. C, $*P<0.0001$ for heterozygous SHR-*Ndufc2*^{em2Mcowi} JD versus SHRSR JD; for SHRSP JD versus SHRSP RD and versus all other samples; $^{\circ}P<0.05$ for SHRSP JD versus heterozygous SHR-*Ndufc2*^{em2Mcowi} JD. F, $*P<0.0001$ for SHRSP JD versus all other samples; $\Delta P<0.01$ for SHRSP JD versus SHRSP RD. H, $\#P<0.001$ and $*P<0.0001$ for SHRSP JD versus all other samples. Het indicates heterozygous; JD, Japanese-style stroke-permissive diet; Nf-kB, nuclear factor kappa B; RD, geular diet; SHRSP, stroke-prone spontaneously hypertensive rat; SHRSR, stroke-resistant SHR; Sod2, superoxide dismutase 2, mitochondrial; wt, wild type.

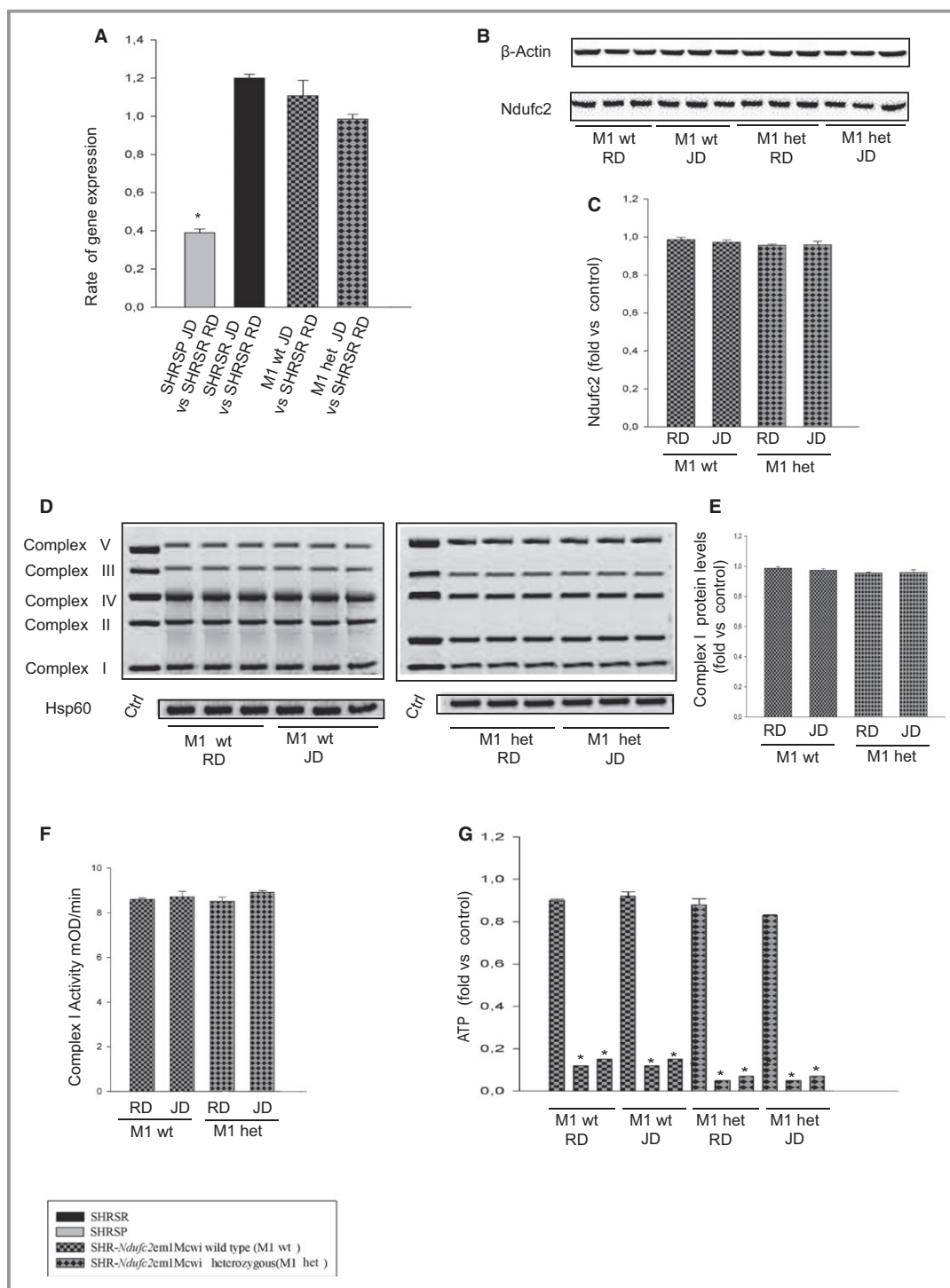


Figure 9. Molecular analyses in the brains of wild-type and heterozygous SHR-*Ndufc2*^{em1Mcwi} at the end of either RD or JD exposure. A, *Ndufc2* expression levels as assessed by RT-PCR. **P*<0.0001 for the indicated ratio (n=4 each). B, Representative western blot of *Ndufc2* in the 2 lines upon either RD or JD with corresponding densitometric analysis (C), number of western blots=3 for each line. D, Representative western blots of BNG (n=3) with corresponding densitometric analysis (E). (F) Complex I activity assessment (n=3 for each group); (G) ATP levels measurement (n=3 for each group). Densitometric values are expressed as means±SD. Ctrl indicates control; het, heterozygous; Hsp60, heat shock protein 60; JD, Japanese-style stroke-permissive diet; RD, geular diet; RT-PCR, reverse-transcriptase polymerase chain reaction; SHRSR, stroke-prone spontaneously hypertensive rat; SHRSR, stroke-resistant SHR; wt, wild type.

levels of proteins underlying necrosis (mitogen-activated protein kinase [MAPK]p38, phosphor-JNK [c-Jun N-terminal kinase], and c-Jun) were significantly augmented ($P<0.001$; $n=3$; Figure 4).

Phenotype and Brain Molecular Analyses of SHR-Derived Lines Carrying *Ndufc2* Deletion

We generated a rat model carrying *Ndufc2* deletion, as described in the Methods section, and characterized its phenotype. No homozygous knockout rat was obtained from either line (SHR-*Ndufc2*^{em1M_{cwi}} [carrying 9-bp gene deletion] and SHR-*Ndufc2*^{em2M_{cwi}} [carrying 107-bp gene deletion]). Furthermore, genotype assessment excluded the presence of homozygous knockout embryos even at early stages of pregnancy, suggesting that *Ndufc2* full deletion is lethal before embryo implantation. Thus, only heterozygous and wild-type rats of the 2 novel rat lines were characterized and compared to the 2 parental strains.

At 6 weeks of age, brains of heterozygous SHR-*Ndufc2*^{em2M_{cwi}} showed a significantly lower amount of *Ndufc2* mRNA ($P<0.01$; $n=6$) and protein expression ($P<0.0001$; $n=6$), as compared to both parental SHRSP and heterozygous SHR-*Ndufc2*^{em1M_{cwi}} (Figure 5A through 5C). A defect in complex 1 assembly ($P<0.0001$; $n=3$) was present only in heterozygous SHR-*Ndufc2*^{em2M_{cwi}} rats, whereas both complex 1 activity and ATP levels were comparable among all strains (Figure 5D through 5I). Level of oxidative stress was also unchanged among strains (data not shown).

JD feeding was started at 6 weeks of age and maintained for 3 months. Systolic blood pressure (SBP), body weight (BW), and proteinuria levels during the dietary treatment are reported in Table 2. SBP levels were comparable among parentals, heterozygous SHR-*Ndufc2*^{em1M_{cwi}} and SHR-*Ndufc2*^{em2M_{cwi}}, and their wild-type controls. Baseline BW was lower in SHRSP ($P<0.001$; $n=18$), as compared to all other lines. Levels of 24-hour proteinuria upon JD feeding increased dramatically in SHRSP and significantly in heterozygous SHR-*Ndufc2*^{em2M_{cwi}}, although to a lesser extent as compared to SHRSP (for all significance values of comparisons at different experimental time points and between strains, see Table 2).

Figure 6 shows results obtained after 4 weeks of either RD or JD feeding in heterozygous and wild-type SHR-*Ndufc2*^{em2M_{cwi}}. At the end of 4 weeks of RD, *Ndufc2* mRNA was significantly decreased in heterozygous SHR-*Ndufc2*^{em2M_{cwi}} as compared to age-matched SHRSP ($P<0.0001$; $n=5$; Figure 6A). JD further reduced *Ndufc2* mRNA ($P<0.0001$; $n=5$; Figure 6A). *Ndufc2* protein expression level was consistent with gene expression data ($P<0.0001$; $n=3$; Figure 6B and 6C). Complex I assembly was significantly reduced in both RD- and JD-fed heterozygous SHR-*Ndufc2*^{em2M_{cwi}} ($n=3$; $P<0.0001$; Figure 6D and 6E). Complex 1 activity ($P<0.0001$; $n=5$),

mitochondrial membrane potential ($P<0.0001$; $n=5$), and ATP levels ($P<0.0001$; $n=5$) were markedly decreased only in JD-fed heterozygous SHR-*Ndufc2*^{em2M_{cwi}} (Figure 6F through 6H). Oxidative stress was also higher in heterozygous, as compared to wild-type, SHR-*Ndufc2*^{em2M_{cwi}} and to parental SHRSP ($P<0.0001$; $n=3$; Figure 7). Nf- κ B protein expression levels were higher ($P<0.01$; $n=3$), whereas those of SOD2 were lower ($P<0.0001$; $n=3$) in heterozygous SHR-*Ndufc2*^{em2M_{cwi}} as compared to SHRSP, similarly to SHRSP (Figure 8). In contrast, all parameters of mitochondrial function and oxidative stress were unchanged in both wild-type and heterozygous SHR-*Ndufc2*^{em1M_{cwi}} (Figures 9 and 10).

Most important, occurrence of stroke reached 100% in SHRSP by 7 weeks of JD and 40% in heterozygous SHR-*Ndufc2*^{em2M_{cwi}} by 3 months of JD feeding. In contrast, stroke did not occur in all other lines (Figure 11A). Comparison of heterozygous SHR-*Ndufc2*^{em2M_{cwi}} versus SHRSP and all SHRSP-derived lines was significantly different ($P<0.002$). Comparison of heterozygous SHR-*Ndufc2*^{em2M_{cwi}} versus SHRSP was significantly different ($P<0.001$).

Analysis of Human Cohort

In the attempt to translate our findings to a clinical setting, we evaluated allelic and genotypic distributions of 3 NDUFC2 tagSNPs in 484 unrelated Caucasian young adults affected by ischemic stroke and in 1165 control subjects. Table 3 shows demographic and clinical characteristics of the study subjects. No statistically significant differences were observed in age and sex between patients and controls. Instead, we observed significant differences in the prevalence of traditional cardiovascular risk factors.

Table 4 shows both genotype distributions and allele frequencies in patients and controls of the 3 studied NDUFC2 SNPs (rs584981, rs641836, and rs11237379). Genotype distributions respected HWE in patients and controls. Prevalence of TT homozygous subjects for rs11237379 SNP was significantly higher in patients than in controls ($P=0.017$; Table 4). The T allele conferred increased occurrence of early-onset ischemic stroke by recessive mode of transmission (OR, 1.39; CI, 1.07–1.80; $P=0.012$). Remarkably, NDUFC2 expression in peripheral blood lymphocytes was significantly and progressively lower in TC/rs11237379 ($n=12$) and in TT/rs11237379 ($n=6$) versus CC/rs11237379 ($n=6$) subjects ($P<0.0001$ for both CT and TT vs CC; Figure 11B).

Even if the difference did not reach statistical significance ($P=0.062$), carriers of the A allele (AA homozygotes and GA heterozygotes) at the rs641836 SNP were more prevalent in patients than in controls (Table 4). We did not observe any difference between patients and controls with regard to the genotype distribution of the rs584981 SNP (Table 4).

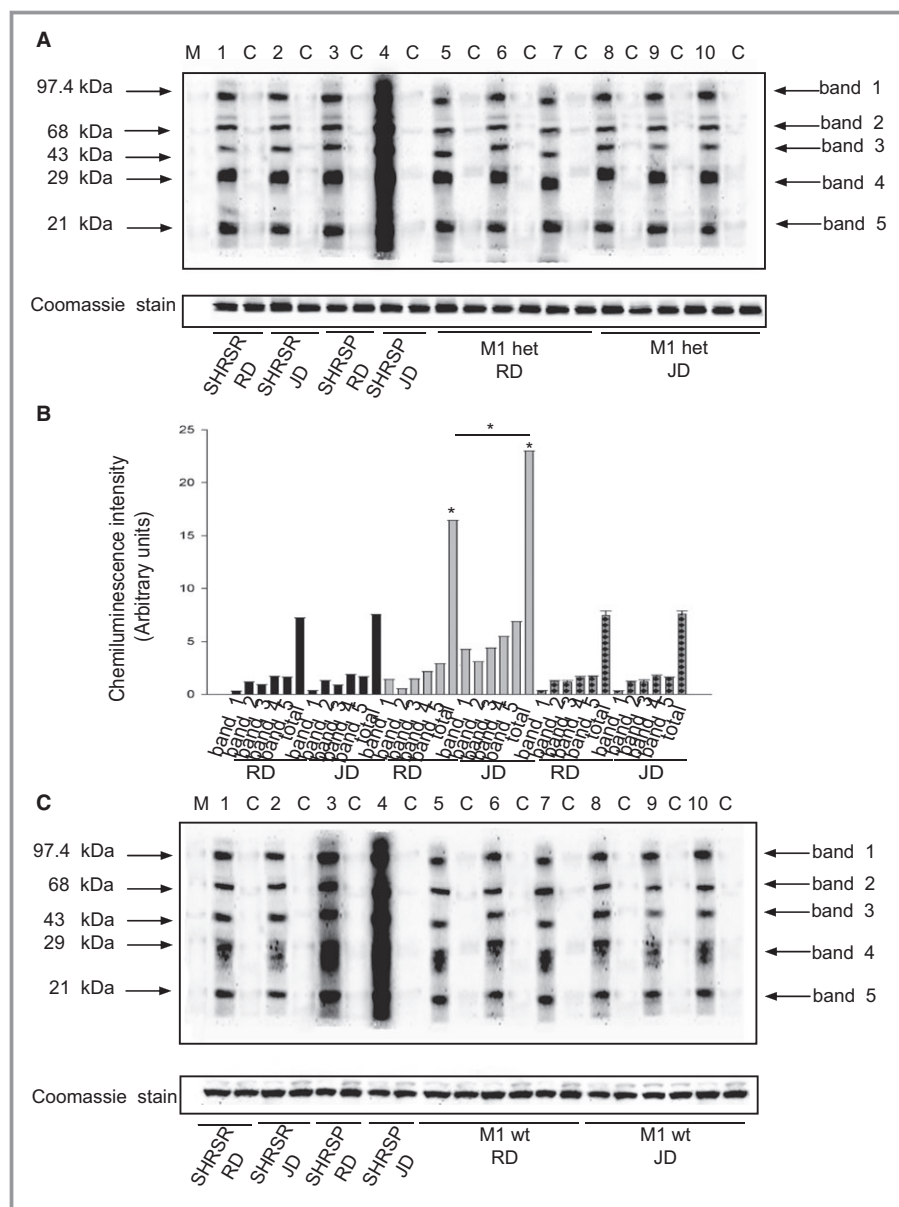


Figure 10. Detection of oxidative stress in the brains of wild-type and heterozygous SHR-*Ndufc2*^{em1Mcwi} at the end of either RD or JD exposure. A, Detection of carbonylated proteins using the Oxyblot Protein Oxidation Detection kit (n=3; Millipore, Milan, Italy) in brains of heterozygous SHR-*Ndufc2*^{em1Mcwi} versus parental lines at the end of 4 weeks of exposure to either RD or JD. B, Corresponding densitometric analysis with values shown as means±SD. **P*<0.0001 for SHRSP JD versus SHRSP RD and for both versus all other samples. Densitometric values are expressed as means±SD. C, Detection of carbonylated proteins using the Oxyblot Protein Oxidation Detection kit (n=3) in brains of wild-type SHR-*Ndufc2*^{em1Mcwi} versus parental lines at the end of 4 weeks of exposure to either RD or JD. D, Corresponding densitometric analysis with values shown as means±SD. **P*<0.0001 for SHRSP JD versus SHRSP RD and for both versus all other samples. Het indicates heterozygous; JD, Japanese-style stroke-permissive diet; RD, geular diet; SHRSP, stroke-prone spontaneously hypertensive rat; SHRSR, stroke-resistant SHR; wt, wild type.

Multivariable logistic regression analysis, with early-onset ischemic stroke as the dependent variable and SNPs and traditional cardiovascular risk factors as independent variables, demonstrated that carrier status of the A allele at

rs641836 SNP or of TT genotype at rs11237379 SNP was associated with increased risk of stroke (OR=1.32 [95% CI 1.03–1.70]; *P*=0.029 and OR=1.39 [95% CI 1.07–1.79]; *P*=0.013, respectively).

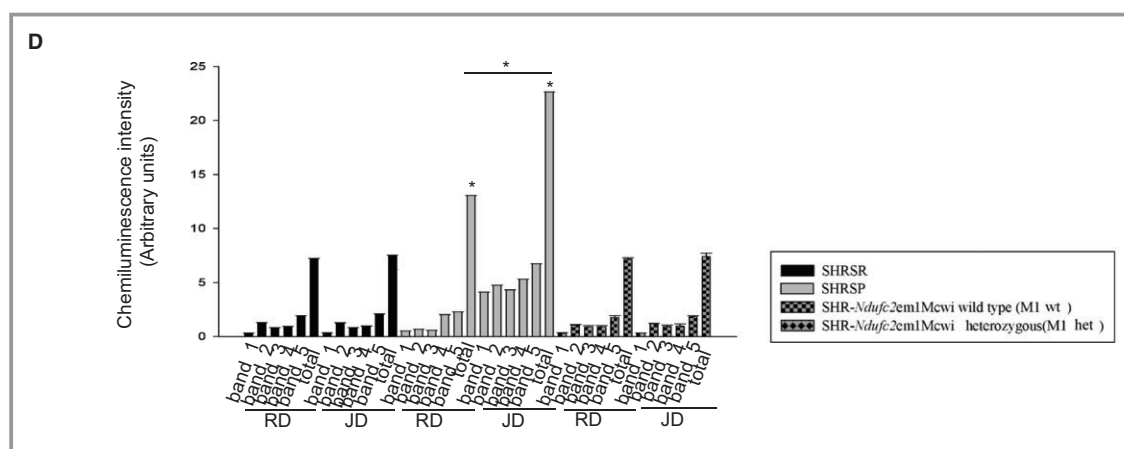


Figure 10. Continued.

At multivariable logistic regression analysis, patients with the combined presence of A rs641836 allele and TT/rs11237379 genotype had the highest early-onset ischemic stroke risk (OR=1.57 [95% CI 1.15–2.14]; $P=0.004$; Figure 11C).

Finally, we observed that the combined presence of A rs641836 allele and TT/rs11237379 genotype was significantly higher in large artery TOAST subtype patients (32.4%) and cardioembolic TOAST subtype patients (40%) with respect to small artery (16%), other cause (19.6%), and undetermined cause (16.0%) TOAST subtypes ($P=0.003$; Figure 12).

Discussion

In this study, we demonstrate that a subunit of NADH dehydrogenase (NADH dehydrogenase [ubiquinone] 1, sub-complex unknown, 2 [*Ndufc2*]), located 8 Mb apart from the Lod score peak of a stroke/QTL on rat chromosome 1, is significantly suppressed by stroke-permissive diet only in brains of SHRSP.

A constellation of cellular abnormalities, including altered complex I assembly and activity, reduced ATP production, increased inflammation, and oxidative stress, was found in brains of SHRSP upon stroke-permissive diet. We demonstrated that *Ndufc2* down-regulation is sufficient to induce several cellular abnormalities and it predisposes to increased occurrence of stroke. By performing in vitro *Ndufc2* silencing in a murine vascular cell line, we provided evidence that *Ndufc2* is a fundamental component of NADH dehydrogenase to allow regular complex I assembly and activity. In fact, its down-regulation led to decreased complex I integrity and function, decreased mitochondrial membrane potential and ATP production, enhanced ROS accumulation, and inflammation. This is consistent with reduced NADH oxidation, which is mediated primarily by complex I. As a consequence, cell viability decreased whereas necrosis significantly increased.

A novel animal model, derived from parental SHRSR, carrying heterozygous deletion of *Ndufc2*, showed increased predisposition to develop both renal and cerebrovascular damage. Hence, we found that brain *Ndufc2* transcription decreased in the presence of 107-bp deletion, as documented by gene expression already at 6 weeks of age. Upon JD, proteinuria preceded stroke occurrence similarly to what happens in SHRSP.³¹ Notably, JD further reduced brain *Ndufc2* expression in this strain and decreased both complex I activity and ATP levels. Interestingly, RD feeding did not compromise complex I activity, despite significantly lower complex I assembly. This evidence suggests that, despite altered complex I assembly in the presence of reduced *Ndufc2* expression, some intrinsic mitochondrial mechanisms can maintain complex I activity and regular ATP levels under RD.³² In addition, this evidence suggest that a high-salt dietary regimen is critical for stroke development in the presence of *Ndufc2* deletion.

Notably, we did not detect any *Ndufc2* mutations in SHRSP, thus suggesting that either epigenetic mechanisms (such as miRNA differential expression) or other gene mutations within the STR1 region may be responsible for *Ndufc2* down-regulation under JD.

Mitochondrion is a major source of energy for cells and its function is fundamental for life of all organisms.³³ NADH dehydrogenase plays important functions in the inner mitochondrial membrane. In fact, it belongs to the oxidative phosphorylation process involved in energy production.³⁴ Electrons are donated from NADH to complex I and then to other components of the chain in order to reduce molecular oxygen (O_2) to form H_2O . Flow of electrons through the OXPHOS chain is accompanied by pumping of protons across the mitochondrial inner membrane, thus creating a trans-membrane electrochemical gradient. Complex I leaks electrons in the intermembrane space to generate superoxide anion (O_2^-). Re-entry of protons into the matrix is used by complex V to synthesize ATP from ADP.³⁴

Complex I is considered the major ROS-generating site within mitochondria. Subunits of complex I are encoded by both nuclear and mitochondria DNA. Mutations in complex I genes lead to increased mitochondrial ROS production.^{33–35} Importantly, the pathological relevance of *NDUFC2* has been recently explored with regard to type 2 diabetes mellitus, providing mechanistic results that are strongly consistent with ours.³⁶ *NDUFC2* appears down-regulated in skeletal muscle cells of subjects affected by insulin resistance³⁷ and is associated with insulin secretion in vivo.³⁸ Furthermore, *NDUFC2* may represent a novel oncogene involved in breast cancer and intestinal adenocarcinoma, predicting poor prognosis.^{39,40} Notably, the pathological relevance of *NDUFC2*/

rs11237379 has been also highlighted through genome-wide association studies in pulmonary disease.^{41,42}

Although it is plausible that mitochondrial dysfunction in general and, particularly, dysfunction of complex I may predispose to cardiovascular diseases and events, no information is available yet with regard to the relationship between *Ndufc2* and cardiovascular disease risk. To our knowledge, the present study is the first demonstration that reduced transcription of *Ndufc2* contributes to stroke.

In summary, *Ndufc2*, encoding a subunit of OXPHOS complex I, mapping close to the lead score peak of a stroke QTL in SHRSP, exerts a key role on stroke susceptibility in this model. A SHRSR-derived line carrying heterozygosity for *Ndufc2* deletion showed increased renal and cerebrovascular phenotypes compared to the resistant strain of origin. The relevance of these experimental findings was extended to a human cohort of early-onset ischemic stroke cases. This population was previously investigated for polymorphisms of candidate genes encoding procoagulant and inflammatory factors and methionine metabolism components: for a polymorphism of type C natriuretic peptide receptor (NPR3) promoter gene and for polymorphisms of chromosome 12p13.^{25–28} In this population, an SNP study revealed a significant association between an intronic marker (rs11237379) of *NDUFC2* and early-onset ischemic stroke. Interestingly, a functional significance of rs11237379 was documented by its direct relationship with gene expression level, suggesting that reduced *NDUFC2* expression may predispose to stroke susceptibility also in humans. Finally,

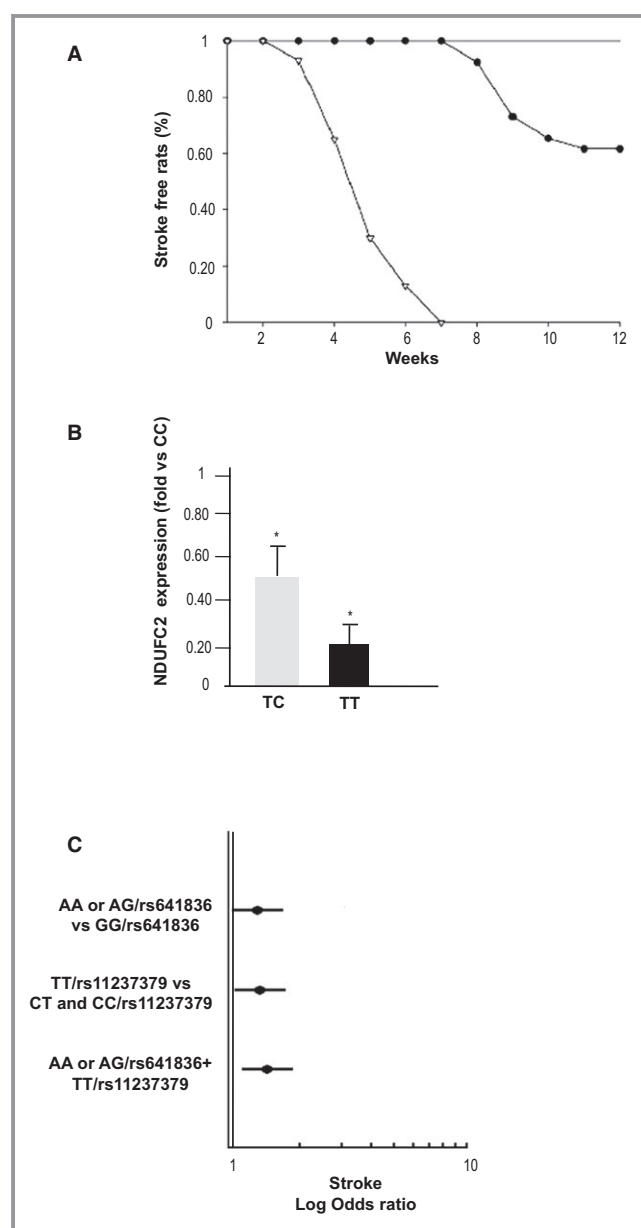


Figure 11. *Ndufc2*-related stroke phenotype in both animal models and in humans. **A**, Stroke occurrence in heterozygous SHR-*Ndufc2*^{em2M_{cwi}} line as compared to SHRSP and to all other lines (SHRSR, wild-type SHR-*Ndufc2*^{em2M_{cwi}}, heterozygous, and wild-type SHR-*Ndufc2*^{em1M_{cwi}}). Comparison of heterozygous SHR-*Ndufc2*^{em2M_{cwi}} versus SHRSR and all SHRSR-derived lines was significantly different ($P<0.002$). Comparison of heterozygous SHR-*Ndufc2*^{em2M_{cwi}} versus SHRSP was significantly different ($P<0.001$). For numbers of animals included in these studies, see Table 1. Symbols: open triangles=SHRSP; closed circles=heterozygous SHR-*Ndufc2*^{em2M_{cwi}}; straight line includes: SHRSR; wild-type and heterozygous SHR-*Ndufc2*^{em1M_{cwi}}, wild-type SHR-*Ndufc2*^{em2M_{cwi}}. **B**, *NDUFC2* expression assessed by RT-PCR in peripheral blood lymphocytes in a cohort of young healthy subjects carrying wild-type ($n=6$), heterozygous ($n=12$), and double-mutant ($n=6$) genotypes for *NDUFC2*/rs11237379. * $P<0.0001$ for both CT and TT versus CC wild-type genotype (t test analysis). Values in the figure are expressed as means \pm SD. **C**, Plot showing the logarithmic ORs for stroke in A allele carriers at rs641836, TT genotype carriers at rs11237379, and A/rs641836+TT/rs11237379 carriers with respect to the other genotypes of *NDUFC2* tagSNPs (at multivariable logistic regression analysis). OD indicates odds ratio; RT-PCR, reverse-transcriptase polymerase chain reaction; SHRSP, stroke-prone spontaneously hypertensive rat; SHRSR, stroke-resistant SHR.

Table 3. Demographic and Clinical Characteristics of Patients With Early-Onset Ischemic Stroke and of Related Controls

	Controls (N=1165)	Stroke Patients (N=484)	P Value
Age median, IQR	42 (32–52)	44 (35–52)	0.078
Sex (male), N (%)	522 (44.8)	222 (45.9)	0.668
Smoking habit, N (%)	297 (25.5)	193 (39.9)	<0.0001
Diabetes, N (%)	4 (0.3)	13 (2.7)	<0.0001
Hypertension, N (%)	85 (7.3)	140 (28.9)	<0.0001
Dyslipidemia, N (%)	39 (3.3)	106 (21.9)	<0.0001

IQR indicates interquartile range.

its pathogenic relevance was enhanced by concomitant presence of a second allele variant at rs641836. Both SNPs may be in linkage disequilibrium with still unknown variants in

Table 4. Genotypes and Alleles Frequency Distribution for rs584981, rs641836, and rs11237379 NDUFC2 Polymorphisms in Early-Onset Ischemic Stroke Patients and in Controls

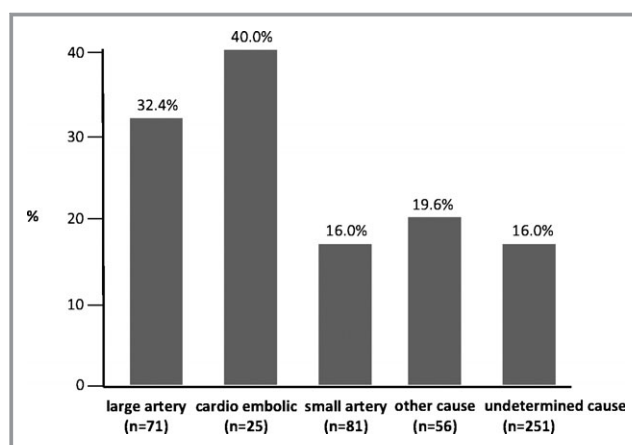
SNP	Controls (n=1165) n (%)	Stroke Patients (n=484) n (%)	P Value
rs584981			
TT	696 (59.7)	294 (60.7)	0.846*
TC	414 (35.5)	170 (35.1)	0.705 [†]
CC	55 (4.7)	20 (4.1)	0.601 [‡]
C frequency	0.225	0.217	
H-W	0.450	0.594	
rs641836			
GG	799 (68.6)	309 (63.8)	0.167*
GA	321 (27.6)	155 (32.0)	0.062 [†]
AA	45 (3.9)	20 (4.1)	0.798 [‡]
A frequency	0.176	0.201	
H-W	2.752	0.010	
rs11237379			
CC	251 (21.5)	94 (19.4)	0.058*
CT	590 (50.6)	227 (46.9)	0.334 [†]
TT	324 (27.8)	163 (33.7)	0.017 [‡]
C frequency	0.469	0.429	
H-W	0.332	0.874	

H-W indicates Hardy-Weinberg; SNP, single-nucleotide polymorphism.

*According to the additive model.

[†]According to the dominant model.

[‡]According to the recessive model.

**Figure 12.** Distribution of the genetic risk condition, derived from combined presence of A/rs641836 allele+TT/rs11237379 genotype for NDUFC2, according to TOAST classification, in a case-control cohort of Italian early-onset ischemic strokes. Prevalence was statistically significantly different (overall $P=0.003$). TOAST indicates Trial of ORG 10172 in Acute Stroke Treatment.

the regulatory or coding regions of the gene. The different distribution of the genetic risk condition, derived from the combined presence of A rs641836 allele and TT/rs11237379 genotype, observed across TOAST stroke subtypes may suggest a prominent role of NDUFC2 in the atherosclerotic and prothrombotic mechanisms underlying stroke.

We are aware that the extension of the experimental results to the human stroke genetic analysis presented in the current work should be considered hypothesis generating rather than definitive, attributable to intrinsic study limitations. In particular, lack of an extensive genotyping for ancestry informative markers in both the autosomal and mitochondrial genomes of our human subjects did not allow a correct population stratification. Additional studies are certainly required to test the hypothesis generated by our experimental data that Ndufc2 may play a contributory role to ischemic stroke predisposition in humans.

Altogether, our novel findings, while pointing to mitochondrial dysfunction as a cause of vascular disease, provide new important insights on the genetic basis of stroke and may have great relevance for prevention and treatment of this common pathological condition.

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Disclosures

None.

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